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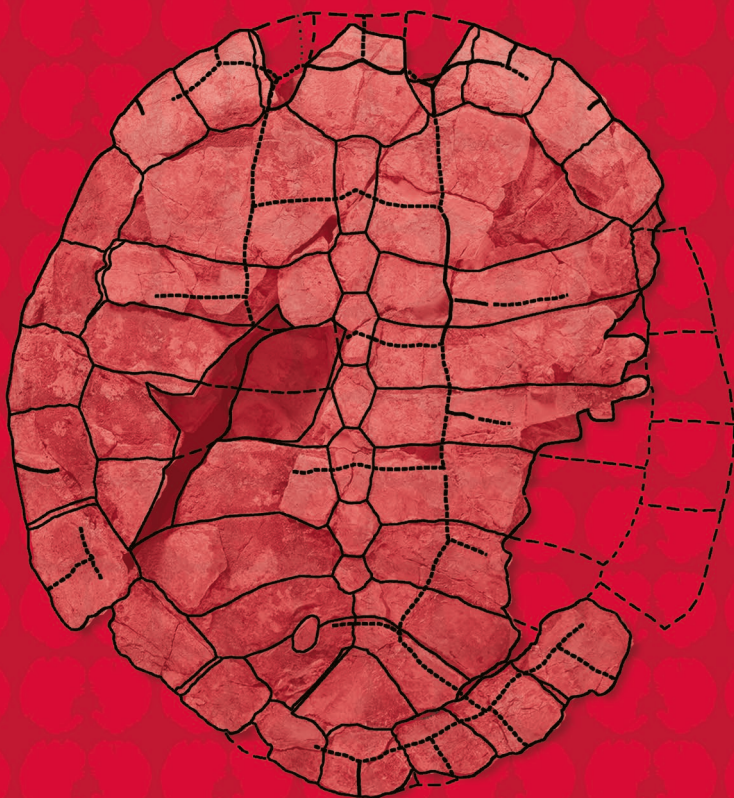
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**First report of a bothremydid turtle,
Sindhochelys ragei n. gen., n. sp.,
from the early Paleocene of Pakistan,
systematic and palaeobiogeographic implications**

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ABSTRACT

We report the discovery of remains of a large chelonian from the base of the early Paleocene Khadro Formation exposed in the Ranikot Fort area (Ranikot Group, Sindh Province, Southern Pakistan). This formation already yielded the snake *Gigantophis* Andrews, 1901, studied by our friend Jean-Claude Rage. The chelonian specimens consist of a large carapace and a shell fragment of Bothremydidae, a family of Gondwanan origin. A new genus and species, *Sindhochelys ragei* n. gen., n. sp. is identified from the first specimen and named in honor of Jean-Claude Rage. It is the first report of a Bothremydidae in Southern Pakistan. Its affinities with Cretaceous and Paleocene representatives of the family are discussed. The association of characters such as the shape of the shell, anterior plastral scute pattern and strongly marked decoration characterize the taxon and, despite some similarities, allows excluding close phylogenetic affinities with Taphrosphyini and *Carteremys* group; other well-documented bothremydids are also excluded. The shell fragment, also strongly decorated, is left undetermined. The discovery of two new littoral bothremydid specimens in the early Paleocene of Pakistan fills a geographic and stratigraphic gap in our knowledge of the family, which is known since the continental early Cretaceous of Africa, diversifying in the world up to the Miocene deposits of the Neotethys. A particular diversification during the Maastrichtian-Paleocene is recognized along the neotethyan coasts, and occasional dispersals across this ocean were possible. *Sindhochelys ragei* n. gen., n. sp. may have colonized the Indian subcontinent by this time, or may represent an older diversification before the Gondwana breakup.

KEY WORDS

Bothremydidae,
Southern Pakistan,
geology,
Pelomedusoides,
Gondwana,
Neotethys,
new genus,
new species.

RÉSUMÉ

Première mention d'une tortue bothrémydide, *Sindhochelys ragei* n. gen., n. sp., du Paléocène inférieur du Pakistan, implications systématiques et paléobiogéographiques.

Nous présentons la découverte de restes de grand chélonien provenant de la base du Paléocène inférieur de la Formation Khadro, affleurant dans la région du Fort Ranikot (Groupe Ranikot, Province du Sindh, Sud du Pakistan). Cette formation a déjà livré le serpent *Gigantophis* Andrews, 1901 étudié par notre ami Jean-Claude Rage. Deux spécimens, une carapace et un autre fragment de tortue Bothremydidae, famille d'origine gondwaniennne, sont décrits. Un nouveau genre *Sindhochelys ragei* n. gen., n. sp. est nommé en l'honneur de Jean-Claude Rage. C'est la première découverte d'un bothrémydide dans le sud-ouest du Pakistan. Les relations avec les représentants crétacés et paléocènes de la famille sont examinées. L'association de caractères comme la forme de la carapace, le pattern des écailles plastrales antérieures et la décoration très accusée le caractérisent et, malgré des similitudes, exclut toute affinité avec les Taphrosphyini et le groupe indien *Carteremys*, les autres bothrémydides bien définis étant aussi exclus. L'autre fragment, également très décoré, reste indéterminé. La découverte de ces deux spécimens dans un environnement littoral du Paléocène inférieur du Pakistan comble une lacune géographique et stratigraphique dans la connaissance des Bothremydidae, connus depuis le Crétacé inférieur continental d'Afrique et s'étant répartis dans le monde jusqu'au Miocène de la Néotéthys. Une importante diversification est reconnue pendant le Maastrichtien et le Paléocène, avec une dispersion le long des côtes néotéthysiennes et peut-être à travers cet océan : *Sindhochelys ragei* n. gen., n. sp. peut avoir colonisé le sous-continent Indien à cette époque, ou résulter d'une ancienne diversification avant la rupture du Gondwana.

MOTS CLÉS

Bothremydidae,
Sud Pakistan,
géologie,
Pelomedusoides,
Gondwana,
Néotéthys,
genre nouveau,
espèce nouvelle.

INTRODUCTION

Bothremydidae is a clade of extinct turtles known from the Cretaceous to Paleogene of African plate, Europe, India, Madagascar, and North and South America. The first record of this group extends back to the early Cretaceous of continental South America and Africa. They represent a monophyletic family of pleurodiran turtles, belonging to the higher group of Pelomedusoides Podocnemidoidea. The family is the sister taxon of the Podocnemididae (Broin in Antunes & Broin 1988; Gaffney *et al.* 2006). Bothremydidae originated on the Gondwana, when Africa, Madagascar, India (including Pakistan) and South America (Brazil) were linked. The oldest representatives of the bothremydids are *Cearachelys* Gaffney,

Campos de Almeida & Hirayama, 2001 from the late Albian of the Chapada do Araripe (Brazil), *Galianemys* Gaffney, Tong & Meylan, 2002 from the late Albian-early Cenomanian of the Hamada du Guir (Kem Kem beds, Morocco) in the upper part of the "Continental Intercalaire supérieur Saharien" (Upper Intercalary Continental of Sahara), and probably the poorly known taxon "*Eusarkia* sp." Bergounioux & Crouzel, 1968, from Tikarkas (Mali) where is recorded the same Albian-early Cenomanian "Continental Intercalaire". Across Sahara, these terrestrial clastics are generally capped by marine beds deposited during the Cenomano-Turonian transgression.

During the Barremian-early Aptian, the family was possibly present in Niger and Brazil where undetermined turtle remains have been reported (Campos de Almeida & Broin 1981).

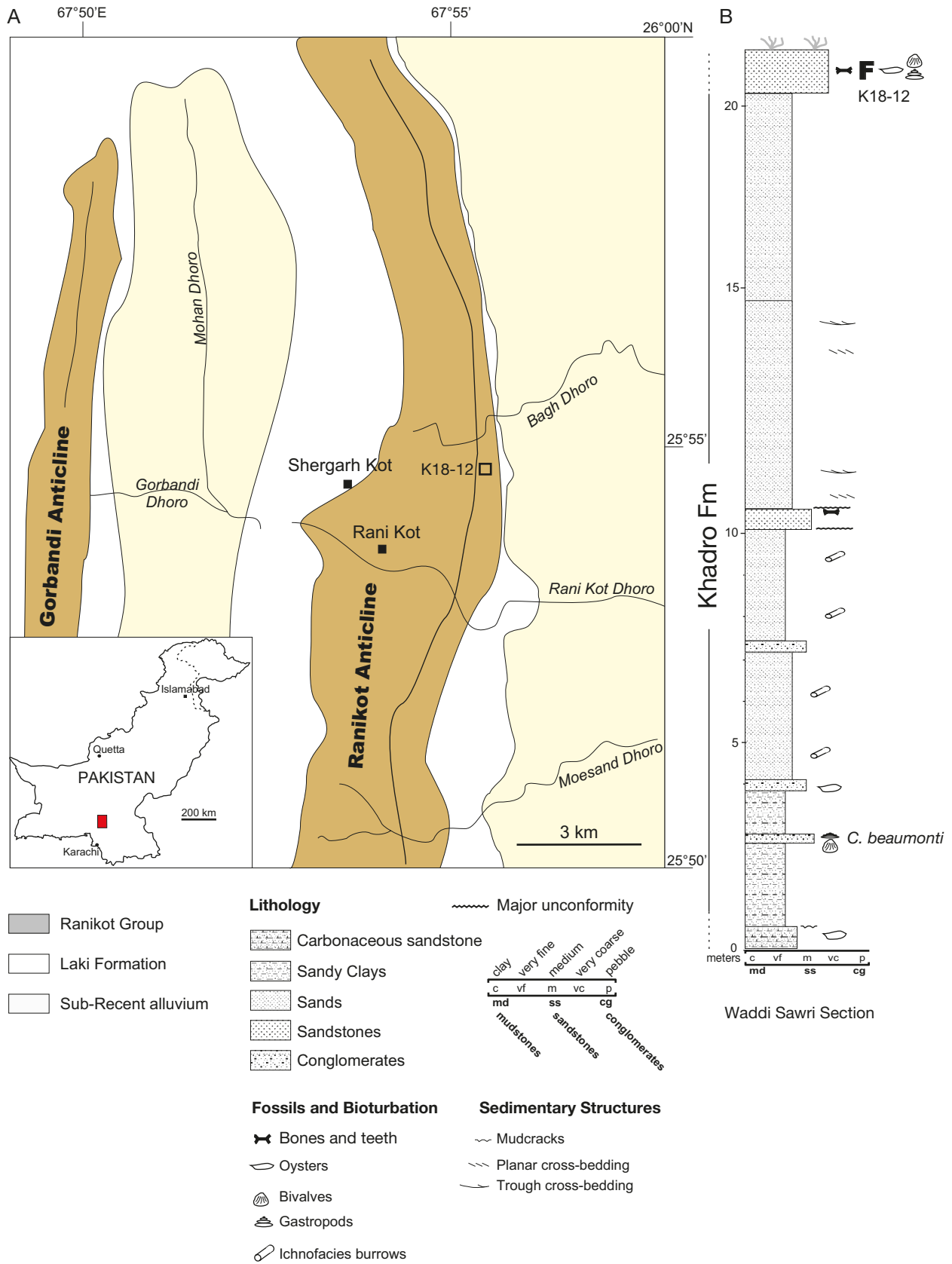


FIG. 1. — Location and stratigraphic position of fossil material: **A**, location of the Laki Range in eastern Sindh Province, southern Pakistan, and simplified geological map of the Ranikot Fort area (adapted from Schelling 1999); **B**, stratigraphic position of the fossil assemblage within the Waddi Sawri section. Note that the base of this section is close to the base of the Khadro Formation; in this area the thickness of the Khadro Formation is c. 80 m, and the fossil locality K18-12 is in the lower Khadro Formation.

TABLE 1. — *Sindhochelys ragei* n. gen., n. sp., holotype CPAG-RANKT-V-3, from the Paleocene of Ranikot, locality K18-12, Southern Pakistan. Shell measurements are in cm.

Body part	Shell measurements
Carapace length – preserved	69.5
Carapace length – estimated	70
Carapace length along curve	72
Carapace width – maximal preserved at peripherals 5/6 junction	54.5
Carapace width – estimated	61
Costal 1, maximal length, left and right	16.5 and 17
Costal 1, maximal width, left and right	13 and 11.5
Plastron, full preserved length	54.7
Plastron, full estimated length	c. 57
Bridge length	c. 20.5
Anterior lobe length up to axillary notch	15
Anterior lobe width at axillary notch	c. 28-29
Posterior lobe length up to inguinal notch	c. 22-23
Posterior lobe width at abdominofemoral sulcus end in inguinal notch	27
Entoplastron, length and width	7.5 and 8
Epiplastral symphysis length	1.7
Mesoplastron, left, length and width	8.7 and 9.8

Moreover, Broin (1980) reported shells of various individuals from the unit 5 of the Elrhaz Formation in Gadoufaoua, Niger (Taquet 1976), which were attributed to the continental freshwater turtle *Platycheiloides* cf. *nyasae* Haughton 1928, a form known from the Cretaceous of Malawi. However, the material from Niger was recognized as clearly distinctive from *P. nyasae*, as noted with M. Cooper (Bulawayo, Zimbabwe) (Lapparent de Broin 2000), and it has been renamed *Francemys gadoufaouaensis* by Pérez-García (2019a). The latter taxon from Gadoufaoua is a Pelomedusoides showing many primitive features and few derived features, such as a marked fine polygonal decoration and the low shell shape with rounded peripheral border, shared with the Bothremydidae. During the Cenomanian, the continental bothremydids became littoral turtles, and widely spread along the coasts of the western Tethys-Atlantic, between Africa, Middle East and Europe, being notably known during the Cenomanian of Palestine and Portugal (Haas 1978a, b; Pérez-García 2016, 2018a; Pérez-García *et al.* 2017a) and up to North America (Joyce *et al.* 2016). They had extensively diversified since the Senonian-Maastrichtian times until their last representatives documented in the Miocene of Oman (Broin 1988; Broin in Antunes & Broin 1988; Roger *et al.* 1994; Lapparent de Broin & Werner 1998; Lapparent de Broin 2000, 2001; Gaffney *et al.* 2006; Lapparent de Broin *et al.* 2018 [online supplementary data]; Lapparent de Broin & Guntupalli 2020). Outside Africa-South America in other Gondwanan continents, the oldest bothremydid is an unnamed taxon from the early Albian-middle Turonian Karai Formation of southeastern India (Ayyasami & Das 1990; Lapparent de Broin & Guntupalli 2020). The family is well documented in the Maastrichtian infratrappean and intertrappean beds of the Deccan volcanic province of India (Carter 1852, Williams 1953; Jain 1977, 1986; Gaffney *et al.* 2001b, 2003, 2006; Lapparent de Broin & Guntupalli 2020) and in Madagascar (Gaffney *et al.* 2006; Gaffney &

Krause 2011; Gaffney *et al.* 2009). Another unnamed taxon is known from the Cenomanian of Madagascar (Lapparent de Broin & Werner 1998; Lapparent de Broin 2000; Lapparent de Broin & Guntupalli 2020). Many turtle taxa are incompletely preserved: some are known by their skull whereas other are so far exclusively documented by their shell (which is the case for the material described here). The result is the studies do not take into account (or only anecdotally) morphological data sets of capital importance (e.g. shell) making biased the attempts of phylogenetic reconstructions (Broin 1988 in Antunes & Broin 1988; Lapparent de Broin & Werner 1998; Gaffney *et al.* 2006) (see Lapparent de Broin & Guntupalli 2020; Lapparent de Broin *et al.* 2020).

Until recently, the reported Pelomedusoides from Pakistan consisted of “*Podocnemis*” *indica* Lydekker, 1885 from the Paleocene of the Salt Range, Northern Pakistan, Eocene undetermined Pelomedusoides (Broin 1987; Smith *et al.* 2016), and early Miocene stereogenyine Podocnemididae (Wood 1970; Prasad 1974; Broin in Antunes & Broin 1988; Gaffney *et al.* 2011; Pérez-García *et al.* 2017b; Ferreira *et al.* 2018). The former was initially reported as “Chelydidae” (corresponding to Podocnemididae), but it has been recently included into the Bothremydidae (Lapparent de Broin & Guntupalli 2020). We show how *Sindhochelys ragei* n. gen., n. sp. cannot be attributed to this form although belonging to the same family. The discovery of two new bothremydid specimens in the Lower Paleocene Khadro Formation (Ranikot Group) of Sindh, Southern Pakistan, fills a gap in our knowledge about the evolution of the family in the Indian Subcontinent. The specimens are notably compared with the ornamented bothremydids including some Paleocene taxa such as *Taphrophys sulcatus* (Leidy, 1856) from the Cretaceous-Danian boundary of New Jersey (United States). New insights into the paleobiogeography of the family are provided.

GEOLOGICAL AND STRATIGRAPHIC SETTING

The term ‘Ranikot Series’ or ‘infra-Nummulitic’, first introduced by Blanford (1876), designates different lithological units sandwiched between the Upper Cretaceous Pab Sandstone and the early Eocene limestone and shales of the Laki Series (‘Kirthar or Lower Nummulitic Group’ of Blanford). Thus, the ‘Ranikot’ is classically considered as Paleocene in age, but the upper and lower boundaries of the Ranikot Group, as those of its three constituent subdivisions (in ascending order, Khadro, Bara, Lakhra Formations), remain poorly constrained chronostratigraphically. The main exposures of the Ranikot Group are in the North-South oriented anticlinal structures forming the Laki Range and extending between Thano Bula Khan in the South and Sehwan Sharif in the North (Fig. 1). In that area, Blanford (1876, 1879) distinguished in ascending order the Cardita beaumonti beds (= *Glyptoactis* (*Baluchicardia*) *beaumonti* (d’Archiac & Haime, 1854)), clastics of the Lower Ranikot, and foraminiferal limestone and sandstone of the Upper Ranikot. The Cardita beaumonti beds is a lithological unit of variable thickness, characterized

by a high concentration of the bivalve *G. (B.) beaumonti*, and these beds have been reported from various spots of eastern Neotethys (Douvillé 1928). Williams (1959) substituted the term Khadro Formation to the *Cardita beaumonti* beds of Blanford (1879) as the lowermost unit of the Ranikot Group and proposed a type section at Bara Nai (c. 20 km north from locality K18-12 that yielded the fossil described here), but the unit has not been properly defined, and the correlation has proven to be difficult laterally. In the Ranikot area, the Khadro Formation consists of olive, reddish to brown sandstone and shale resting unconformably on the Khaskheli altered basalts (Shah 2009; Agheem *et al.* 2011), the latter overlying the Campano-Maastrichtian Pab Formation.

Deposition of the Khadro Formation is interpreted to have occurred along the approximately North-South trending passive margin of eastern Neotethys, and the lithology and fossil content (Rage *et al.* 2014) suggest estuarine-coastal to inner shallow platform environments, which conforms to the known ecology of Bothremydidae.

Uplift and compression have been episodic since the Paleocene, but the main phase of compression, folding and uplift was during the Pliocene to the present (Schelling 1999).

The fossil specimens were collected on surface of a large platform of indurate sandstone (locality K18-12), which has also yielded the giant snake *Gigantophis* (Rage *et al.* 2014), crocodile remains, oysters, bivalves (?*Crassatella* sp.), and gastropods (*Haustator* sp.), suggesting a coastal depositional environment. The stratigraphic position of the locality K18-12 is indicated on the Waddi Sawri section (Fig. 1), which exposes the lower part of the Khadro Formation, including the *Cardita beaumonti* beds. The fossiliferous horizon is stratigraphically situated about 20 meters above the horizon of carbonaceous green sandstone showing the higher concentration of complete bivalves of *G. (B.) beaumonti*. The entire and measured Waddi Sawri section indicates that the thickness of the Khadro Formation does not exceed 80 meters in the Ranikot area. The complete carapace and plastron of turtle described here were recovered from the lower part of the Khadro Formation, just above the famous *Cardita beaumonti* beds (Blanford 1879). These beds lying near the base of the Khadro Formation indicate an early Paleocene (Danian) age (Cheema *et al.* 1977; Shah 2009). The Danian age of the Khadro Formation is supported by bivalves (Eames 1952) and foraminiferals (Nagappa 1959) assemblages. Thus, the bothremyd turtle described here is considered as early Danian in age.

MATERIAL AND METHODS

The shell and other fragments of turtles described here were collected during the 2012 field season from the same stratigraphic interval that yielded the giant snake *Gigantophis* (Rage *et al.* 2014). Because of its weight, the fossil shell was taken out from the sandstone in three blocks (Fig. 2) in order to make smaller plaster jackets to facilitate its long transportation to the field vehicle. The fossil shell was reassembled and meticulously prepared in Paris by Philippe Richir (CR2P), and Yohan Després (CR2P) made a resin jacket to facilitate

its manipulation during detailed fossil study. In course of the preparation, many, marine and dwellers bivalves (Veneridae), have been found in the sandstone. They are articulated anatomical connection and probably still in live position (Fig. 2A1). This is consistent with the inferred littoral ecology of bothremydids, most of the time. Taphonomically, the occurrence of venerids in life position inside the carapace indicates that the turtle was buried in low energy littoral environment. In the terminology, the plates overlapping the thoracic ribs are named costals and the scutes which cover them are named pleurals. The measurements are given in cm in Table 1.

ABBREVIATIONS

Institutions

AMNH	American Museum of Natural History, New York;
BRGM	Bureau de Recherches géologiques et minières;
CPAG	Centre for Pure and Applied Geology, University of Sindh, Jamshoro, Pakistan;
DNPM	Departamento Nacional da Produção Mineral, Rio de Janeiro;
ENSMP	École nationale supérieure des Mines, Paris;
GSI	Geological Survey of India, Kolkata;
MFL	Main Fossiliferous Layer of the Hornerstown Formation;
MHNM	Musée d'Histoire naturelle, Marseille;
MNHN.F	Muséum national d'Histoire naturelle, collection de Paléontologie, Paris;
MRAC	Musée royal de l'Afrique centrale, Tervuren;
NJ	New Jersey;
PU	Princeton University.

SYSTEMATIC PALEONTOLOGY.

Order TESTUDINES Linnaeus, 1758
 Suborder PLEURODIRA Cope, 1864
 Hyperfamily PELOMEDUSOIDES Cope, 1868
 Superfamily PODOCNEMIDOIDEA Cope, 1868
 Family BOTHREMYDIDAE BAUR, 1891

Genus *Sindhochelys* n. gen.

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TYPE SPECIES AND ONLY SPECIES OF THE GENUS. — *Sindhochelys ragei* n. sp.

DIAGNOSIS. — Same as the type species of the genus.

ETYMOLOGY. — From Sindh, the province of discovery in Pakistan.

Sindhochelys ragei n. gen., n. sp.
 (Figs 2-9)

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TYPE MATERIAL. — Holotype, a shell, CPAG-RANKT-V-3. The final repository of the fossil material reported here is the Centre for Pure and Applied Geology (CPAG), University of Sindh, Jamshoro, Pakistan. High-resolution casts are housed at the Muséum national d'Histoire naturelle, Paris.



FIG. 2. — Turtle shell, CPAG-RANKT-V-3, holotype of *Sindhochelys ragei* n. gen., n. sp., *in situ*, during and after excavation at the locality K18-12: **A**, carapace in posterior view; **A1**, close-up of a lateral border showing the presence of a venerid bivalve in life position inside the carapace; **B**, carapace in dorsal view after excavation and split into three blocks. Scale bar: 30 cm.

DIAGNOSIS. — For monotypic genus and species. A bothremydid turtle defined by the ornamentation of the shell consisting of raised well protruding polygons but, unlike other ornamented forms such as Taphrosphyini and *Carteremys leithii*, in having conjoined rather larger irregular polygons without a clear net of granulations, polygons clearly but irregularly protruding and either elongated or regularly polygonal, or open in dichotomic sulci at places. Sharing with *Elochelys perfecta*, *Carteremys* group, “*Carteremys*” *pisdurensis*, and Taphrosphyini the generalized plastral anterior lobe scute pattern of a long intergular, separating narrower and shorter gulars, and widely reaching the pectorals, intergular also separating the thus narrowed humerals; but: humero-intergularo-pectoral sulcus crossing the entoplastron at the 2nd anterior third of its length, its humero-pectoral part posterior to the epi-hyo-plastral suture in its

median width, but close to it and laterally joining it, contrarily to *E. perfecta* where it is completely anterior to the suture, contrarily to “*C.*” *pisdurensis* where it is medially anterior to the suture and lateroposteriorly inclined much posteriorly to the suture, and contrarily to Taphrosphyini and the *Carteremys* group where it does not cross the suture; intergularo-pectoral sulcus relatively transverse on the entoplastron, as in the *Carteremys* group, not being medially inclined in a marked V contrarily to *E. perfecta*, “*C.*” *pisdurensis* and Taphrosphyini. Bothremydid carapace low with well-inclined bridges but different from other forms, being wider at the suture of peripherals 6-7; displaying a combination of other characters which are distributed in mosaic in the bothremydid groups such as: carapace anteriorly relatively elongated (with regard to primitive forms), at costals 1, neural 1 and anterior peripherals, but moderately, and

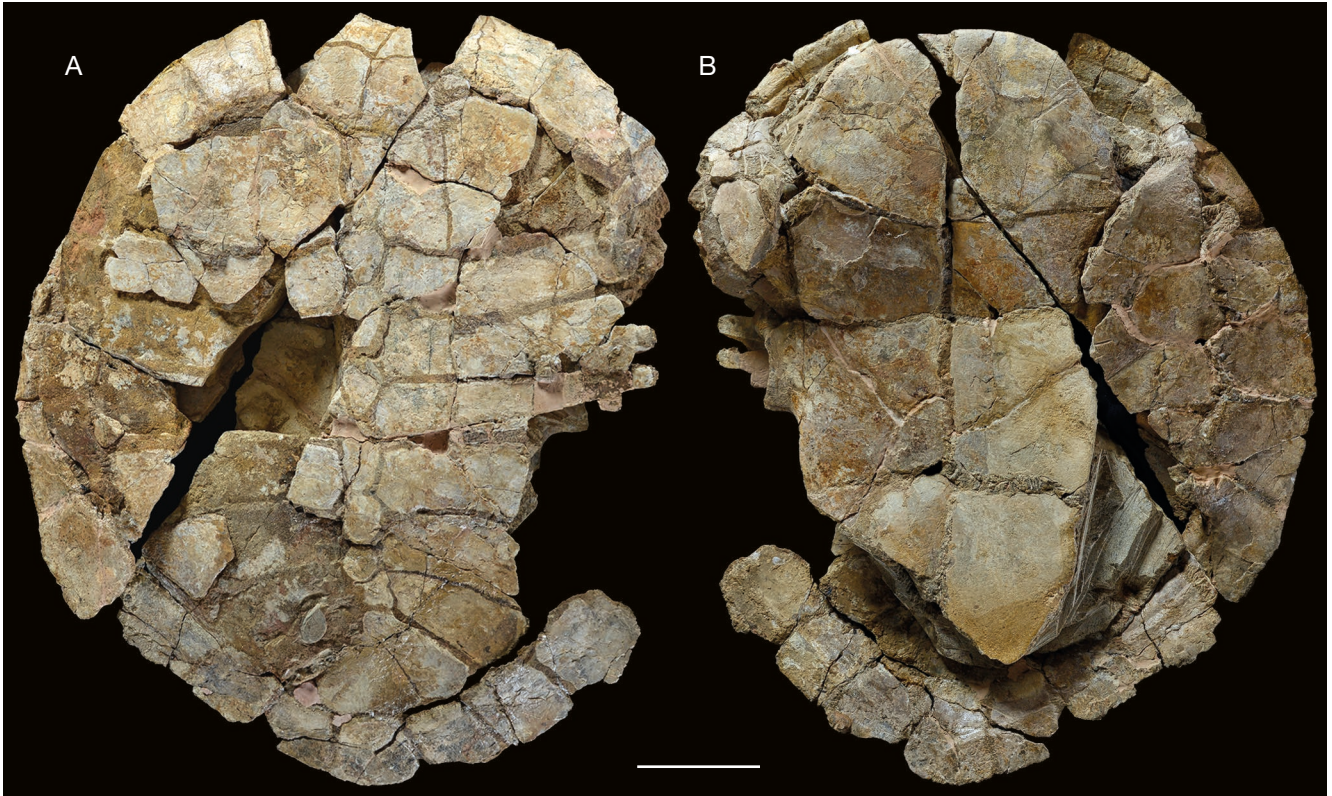


FIG. 3. — *Sindhochelys ragei* n. gen., n. sp., holotype CPAG-RANKT-V-3, from the Paleocene of Ranikot, locality K18-12, Southern Pakistan: **A, B**, shell photographs in dorsal (**A**) and ventral (**B**) views. Scale bar: 10 cm.

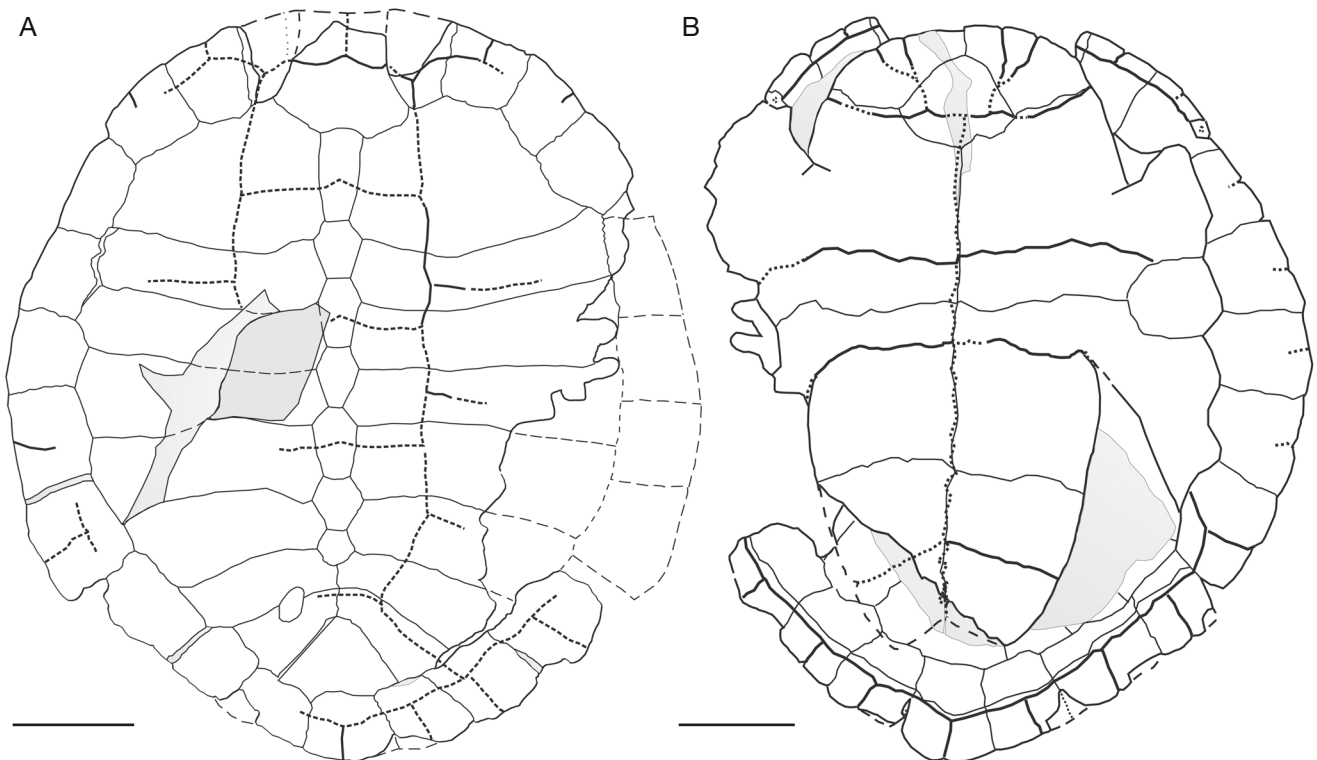


FIG. 4. — *Sindhochelys ragei* n. gen., n. sp., holotype CPAG-RANKT-V-3, from the Paleocene of Ranikot, locality K18-12, Southern Pakistan: **A, B**, shell drawings, in dorsal (**A**) and ventral (**B**) views. Scale bars: 10 cm.

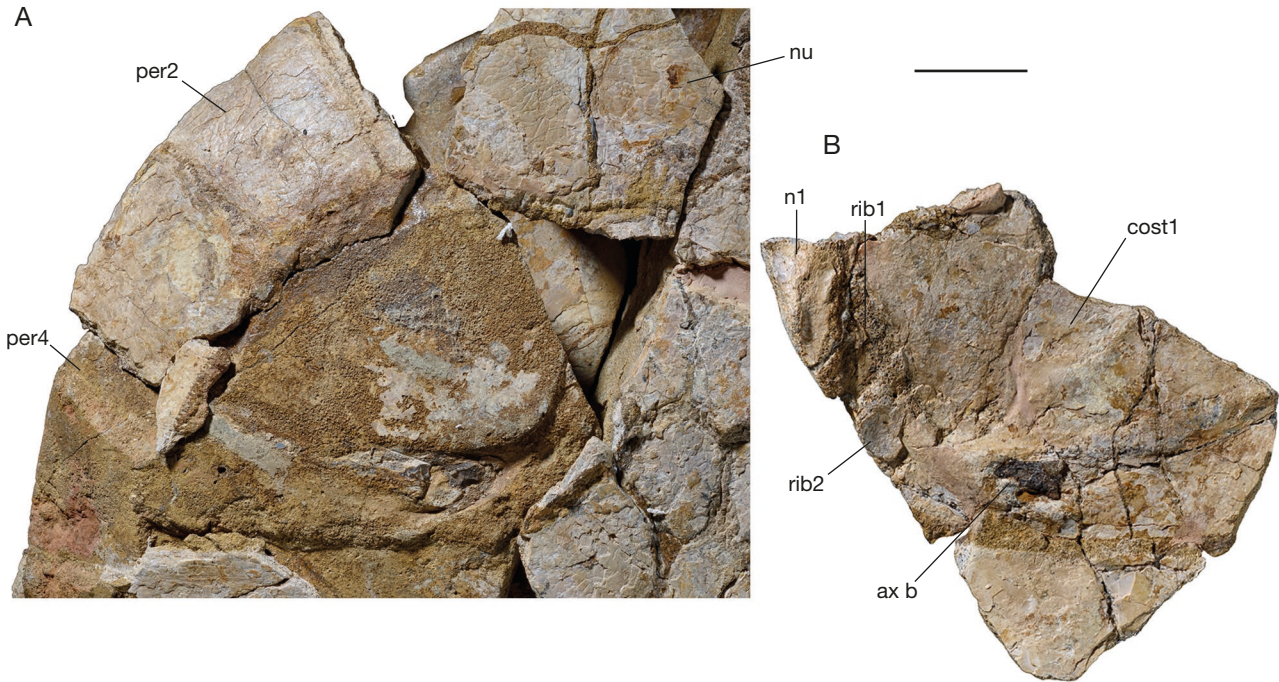


FIG. 5. — *Sindhochelys ragei* n. gen., n. sp., holotype CPAG-RANKT-V-3, from the Paleocene of Ranikot, locality K18-12, Southern Pakistan: **A**, anterolateral part of the carapace after removal of a part of the costal 1 with part of the neural 1 and the costal 2, to show the imprints of the thoracic ribs 1 and 2, and axillary buttress, in dorsal view; **B**, same removed part in ventral view. Abbreviations: **ax b**, axillary buttress; **cost 1**, costal 1; **n1**, neural 1; **nu**, nuchal; **per 2**, peripheral 2; **rib 1**, **rib 2**, medial part of thoracic ribs 1 and 2. Scale bar: 4 cm.

with an anterior lateral convex contour and without medial nuchal projection; rounded marginal free border of the anterior peripherals; posterior contour slightly protruding towards pygal in a wide convex V; seven relatively narrow neurals; anterior and posterior (overall) plastral lobes not reaching the carapace borders in length; short trapezoid anterior lobe with lateral borders converging only a little towards the very gently convex transverse front, wider at base than the posterior lobe that is long; axillary buttresses laterally beginning at suture peripherals 3 – 4, not particularly thinned, medially overlapping in a curve two thirds of the thoracic rib 2 and medially ending in a rounded extremity on the rib 2 and, contrarily to *Taphrosphys sulcatus*, rib 2 not close to the costals 1-2 suture towards the median part; rib 1 much anteriorly standing up close to the suture between costals 1 and 2; bridge (between axillary and inguinal notches bottom) longer than the anterior lobe and shorter than the posterior one, with upwards inclined lateral processes, massively shaped around the large (nearly as long as wide) mesoplastra and beside the wide bridge peripherals; posterior lobe with gently convex lateral borders little posteriorly converging. Long marginals 1 overlapping at least the anterior mid-length of the nuchal, and marginals 2 overlap laterally much shortening, in convex curve from peripherals 1 to peripherals 2, this short marginal overlap extending at least to the fourth peripherals (not preserved in situ on the specimen up to the peripherals 8), and with a short ventral marginal overlap; dorsally these short lateroanterior marginals correspond to long pleurals 1, longer in relation to the following pleurals 2 to 4; vertebral 1 a little wider than the nuchal, 2nd to 4th as wide; possibly 4th the longer; and 5th the shorter, but posteriorly the wider. As in various other bothremydids, pectoroabdominal sulcus convex and slightly anterior to the mesoplastra, and curving towards them at its lateral extremity, and not posterior on the hyoplastron all along, contrarily to *Taphrosphyini*; femorals longer than pectorals, both long in relation to abdominals and anals.

ETYMOLOGY. — From *chelys*, turtle in Latin. Species: in honor to our friend and colleague Jean-Claude Rage.

HORIZON AND TYPE LOCALITY. — Lower part of the Khadro Formation, Ranikot Group, early Paleocene (probably early Danian in age), type locality K18-12, in the proximity of Ranikot Fort, Jamshoro District, Sindh Province, Southern Pakistan (Fig. 1).

DESCRIPTION (Figs 2-10)

Preservation

The shell has been flattened during the fossilization, and the plastron is fractured at bridges and slightly pushed into the carapace. The carapace is therefore a little wider than when the individual was alive. A great part of the carapace was imbedded in a hard matrix, which could not be removed (Fig. 2), and a part of the plates were dislocated inside the whole assemblage. After the preparation, the elements of broken plates were not in contact, and they are preserved separately; some parts remained in the hard matrix. The photograph (Fig. 3) shows the preserved prepared part with plates in place or in imprints allowing a drawing of the full carapace and plastron (Fig. 4).

Dimensions (Table 1)

The shell is large (around 70 cm long) and the length/width ratio on (ca. 85 to 90 %) shows it was relatively high for the family, even taking into account the post-mortem flattening. The shell was clearly wider than in the more primitive Pelomedusoides and podocnemidoid forms of Brazil and Morocco. It is neither the smallest nor the largest known bothremyd fossil shell. The oldest and primitive Brazilian bothremyd *Cearachelys* has a shell length of c. 15 cm (Gaffney *et al.* 2006). Whereas some Maastrichtian shells from Niger have a length of 120 to 150 cm (Lapparent de Broin *et al.* 2020). Although much flattened post-mortem, the shell was however rather

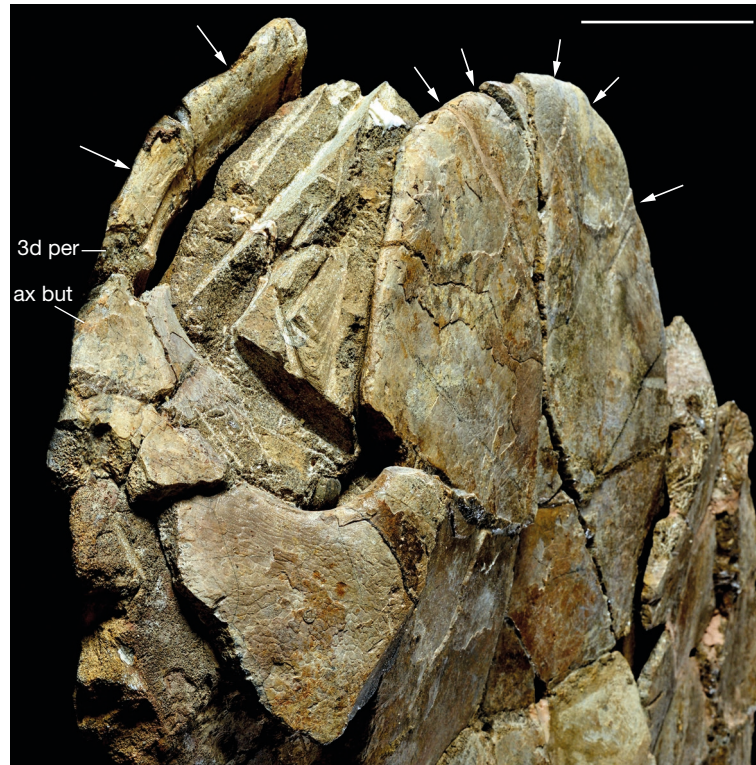


FIG. 6. — *Sindhochelys ragei* n. gen., n. sp., holotype CPAG-RANKT-V-3, from the Paleocene of Ranikot, locality K18-12, Southern Pakistan: left anterior part of the shell in oblique posterolateral view showing the delimitation of marginals 2 and 3, and of plastral scutes by the indentation at the end of sulci border, and the ventral bulges of the scutes. Abbreviations: **ax b**, axillary buttress; **3d per**, third peripheral. The arrows indicate ends of sulci at the border of the peripherals and anterior plastral lobe. Scale bar: 5 cm.

low and gently rounded dorsally, without dorsal carina, and with well inclined bridges.

Carapace (Figs 2-9)

The maximum width is at the suture between the peripherals 6 and 7 (Figs 3A; 4A). There are no parallel lateral borders on a short length between the peripherals 4 to 8, unlike most bothremydids: the contour widens from peripherals 2 to 6 and narrows from peripherals 6-7 contact up to the pygal. Anteriorly, the outline of the carapace is semicircular, being just straight at the nuchal area border (Fig. 4A). Posteriorly, the carapace reduces in width with a medial posterior protrusion, forming a wide V. Thus, the carapace was not ovoid in shape, and its anterior part was not anteriorly longer and narrower, in relation to the less long posterior part. The anterior border of the nuchal 1 and most of the border of peripherals 1 is not preserved, but the contour may be reconstructed as transversally straight at the nuchal (Fig. 4A) and at medial part of peripherals 1. Indeed, this part does not seem to have been much projected because of the very moderately inclined border of peripherals 2 and 3; this character differs from the condition seen in some taphrosphyine (*sensu* Lapparent de Broin & Guntupalli 2020) bothremydids such as “*Gafschelys neurriregularis*” Bergounioux, 1952 from the Ypresian of Gafsa (Tunisia) (Bergounioux 1956, pl. 10). Besides, the median carapace part has no parallel lateral borders, and the posterior part is not rounded as in some bothremydids such as *Elochelys perfecta* Nopcsa, 1931a (Nopcsa 1931b) from the Campanian

of Faveau-Valdonne (France). The anterior part of the shell is elongated in relation to the primitive condition seen in the oldest bothremydids such as *Cearachelys* (Gaffney *et al.* 2006). The maximal length of the costal 1 (at the suture of peripherals 2 and 3) is here as large as the length of the part including the costals 2, 3 and 4 at their mid width (measured following the surface curve). In agreement, the nuchal and peripherals 1 and 2 are also elongated. Although its anterior border is missing, it is clear the nuchal was elongated, but shorter than the costals 1 (Fig. 4A). The free border of peripherals 2 and 3 is not acute; it is rounded as in some of the indetermined Bothremydidae from the Maastrichtian of Upparhatti (India) (Lapparent de Broin & Guntupalli 2020: fig. 7); it is ventrally rounded with a short marginal scute overlap (Figs 3B; 6). The costal 1 is united by concave sutures: 1) anterolaterally to the posterolateral nuchal border and to peripherals 1 to 4, and medially to the neural 1; 2) and by straight sutures to the anterolateral border of the neural 2 and with the costal 2. The first peripheral is the narrowest of all the series, and it is overall narrow posteriorly, at its suture with the costal 1. There are seven neurals. The neural 1 is narrow and elongated in agreement with the costal 1. It is quadrangular, the longer of the series (in agreement with the costal 1 great length). Neurals 2 to 5 (hexagonal with short anterolateral sides) are narrow, and an individual variability in their shape and relative length is observed: 1) they have a narrow and nearly equidimensional contact between them but a narrower contact is situated between the neurals 4 and 5; and 2) they have a slightly different

length, and a variable maximal width (at the junction of successive costals). Hexagonal neural 6 is shorter than neurals 2 to 5, and pentagonal neural 7 is even shorter, being the shortest in the series: both 6th and 7th have lateral sides which are as long anteriorly and posteriorly. The costals 7 partly posteriorly meet in the mid-line behind neural 7; the costals 8 are fully meeting medially. Costals 2 to 4 are approximately as long laterally as medially, costals 5 are laterally longer, and costals 6 to 8 are laterally longer and less wide (overall the 8th), accompanying the posterior carapace protrusion. The neurals 2 to 5 appear long because they are relatively narrow for their length, particularly the neurals 3 and 4, and evidently their length is in agreement with the costal length, i.e. they are not particularly long in the carapace. The suprapygal is pentagonal as usual in the family when the posterior costals 8 meet in the midline and the pygal is quadrangular, longer than wide in dorsal views (Figs 3A; 9C). In the dorsal views, the posteromedially pointed posterior border of the carapace appears shorter than the anterior one because the carapace is more inclined than the anterior border. On the left, part of the costal 6 and costals 7 and 8 are seen only by their impression: the transection of the iliac bone appears as an oval part in the matrix, between the 7th and 8th plates impression (Figs 3A; 9C), as usual below its suture with these plates.

It has been possible to separate a fragment with a good part of the neural 1, the left costal 1 and a triangular medial part of the costal 2, which allows to see its ventral aspect (Fig. 5B) and the corresponding impression of the carapace in dorsal view (Fig. 5A). The thoracic rib 2 is extended, below the costal 1 surface, medially from the interval between the neurals 1 and 2 towards laterally the peripheral 4; all along its course, it is rather close to the suture with the costal 2 but not very close, and not linked to it as in some forms. The thoracic rib 1 first extends (from laterally to medially) below the costal 1 along the medial third of the rib 2. Close to the medial border of the costal 1, rib 1 abruptly curves to extend anteromedially towards the level of the mid-length of the neural 1, where it sutured with the apophysis (not preserved) of the thoracic vertebra. The axillary buttress extends medially upon the thoracic rib 2 in a gentle curve, from the suture between the peripherals 3 and 4, and it covers the rib 2 on the two lateral thirds (Fig. 5A, B), making a rounded bulge ventrally (Fig. 5B). It medially ends in a rounded scar which is not dilated. In their median part, the axillary buttress and rib 2 are not as close to the suture between the costals 1 and 2 than in taphrosphyine specimens (Lapparent de Broin & Werner 1998).

Plastron (Figs 3B; 4B; 6; 7)

It is nearly complete, missing most of the right xiphiplastron and the end of the left xiphiplastron, close to the anal notch, at the insertion of the ischion. The pelvis is concealed by the matrix and the shape of the pubis and ischiatic scars are unknown. However, in posterior view of the shell, the ischiatic contact with the xiphiplastron appears as making a transverse contact: there is no indication of the rounded scar of *Taphrosphys sulcatus* (Leidy, 1856) and other members of the taphrosphyine tribe (Gaffney 1975: fig. 11; Gaffney *et al.* 2006) (this tribe corresponding here to the *Taphrosphys* group

of Broin in Antunes & Broin 1988, i.e. *Taphrosphyini sensu* Lapparent de Broin & Guntupalli 2020). The axillary notches are slightly deformed, better preserved on the right side. The bottom of the right inguinal notch is preserved, and the left one is partly cut by a crack; but its bottom position is indicated by the lateral end of the abdominofemoral sulcus on the hypoplastron, beside the crack, and conform to the right preserved side. The bridge length between the notches bottom was long, reaching approximately 36% of the full length of the plastron. The axillary buttress reaches the boundary between the peripherals 3 and 4 (Fig. 6), and the inguinal buttress reaches $\frac{2}{3}$ rd of the peripheral 8 (Fig. 3B, left side).

The anterior lobe is wide so that it laterally covers rather well the axillary notches space, much occupying the anterior carapace opening in ventral view. It is trapezoidal with a wide transverse anterior border, barely convex, wide at its base and not reaching the anterior nuchal border. The epiplastral symphysis is short and the entoplastron is large in the anterior lobe. It is widely pyriform and its posterior extremity is posterior to the transverse line passing at the bottom of the axillary notches (Fig. 3B, right side). The bridge length between the axillary and inguinal notches is nearly as long as the posterior lobe. The bridge presents the great oblique surfaces of the lateral processes between the main plastral body and the corresponding ventral border of the wide peripherals. The large lateral mesoplastra are hexagonal, a little wider than long, their length representing about one third of the lateral bridge length. The posterior lobe is slightly less wide at its base than the anterior one. Its lateral borders, barely rounded just posterior to the inguinal notch bottom, are gently converging posteriorly towards the mid-line. The posterior lobe does not cover as much the inguinal notches space than in some other bothremydids, but more than in the primitive condition, in which lateral borders are straighter and more posteriorly converging (e.g., *Francemys* Pérez-García, 2019 [Pérez-García 2019a] and *Cearachebys* Gaffney, Campos de Almeida & Hirayama, 2001 [Gaffney *et al.* 2001a]). It is likely that the anal notch was wide and short (doing a short and wide triangle).

The plastral plate length formula is: anterior lobe < bridge < posterior lobe length.

Shell scutes

An important part of the sulci is not preserved. Some are clearly visible, other are estimated (Fig. 4[dotted lines]). When not visible, their lateral extremity on the border of the element is positioned by the corresponding small incurved border of the plates, and by the sulcus continuation on the inferior face of the plates: this is seen by the border of marginals on the peripheral plates and by the border of plastral scutes on the epi-hyoplastra (Figs 3B; 4B; 6).

The carapace (Figs 3; 4) has long marginals 1 on the nuchal. These scutes are estimated as overlapping about the nuchal anterior mid-length (Fig. 4A). Dorsally, the sulci between the two marginals 1 and the vertebral 1, corresponding to the posterior borders of the marginals 1, are slightly concave. On the nuchal, the curve of the sulcus between the vertebral 1,

and marginal 1 and the sulcus between the marginal 2 and the vertebral 1 on the peripheral 1 helps identifying the possible position of the missing sulcus between the marginals 1 and 2: this was close to the nuchal-peripheral 1 suture (Fig. 4A [dotted line left side]). Each marginal 1 had to be a rectangular scute and barely wider than long. The marginal 2 was wide, long medially but laterally shorter, from the indicated right curve on the peripheral 1. It was followed by a short marginal 3. The following marginal sulci are mostly missing but from the 7th marginal, on the peripherals 6 and 7, an overlap elongation is estimated, thanks to the preservation of parts of some sulci: as a whole, the marginal dorsal overlap seems to widen at the bridge and slightly narrow down after the bridge (from the peripherals 9 up to the pygal). This medioposterior marginal overlap is longer than that on the anterior peripherals. In conformity, ventrally, the marginals 8-9 sulcus, which is preserved on the left peripheral 8, shows that the ventral marginal overlap is wide at this bridge 8th peripheral, and then the ventral overlap is slightly reduced in length posteriorly. The anterior ventral carapace face indicates a short scute overlap and a rounded thick border (Figs 3B; 6). The ventral scuteskin boundary is marked: anteriorly to the bridge by a thin groove along a low bulge, and posteriorly to the bridge by a protruding rounded half-bulge. The vertebral 1 was wider than the nuchal (thanks to the preserved beginning of the sulcus on the right peripheral 1), the second (its right border being preserved) and third are estimated as equally wide or barely wider than the first, and the fourth seems to be posteriorly narrowed and in contact with the fifth vertebral on the costals 8. If the vertebrae are correctly reconstituted, the longer vertebral is the 4th, as long as wide, while the 1st, the 2nd and 3rd are wider than long. By the preserved parts of sulci, the vertebrae 1 to 4 seem to be as wide all along, and they are not narrowed at their mutual contact, contrarily to the 4th at its contact with the 5th. From the width of the posterior costals and their contact with the peripherals, it seems the vertebral 5 was posteriorly wider, on the peripheral 10, wider than the other vertebrae. Because of the shorter marginals 2 and 3 in *Sindhochelys ragei* n. gen., n. sp., the pleural scute 1 was the longest of the series and more than in many derived forms, those which also have an elongated anterior border.

The plastral scute sulci are mostly preserved (Figs 3B; 4B; 6). The anterior lobe scute pattern of *S. ragei* n. gen., n. sp. is estimated as a derived complex of scute arrangement. It is found in several taxa of several different podocnemidoid groups, but the details in the sulci outlines and scute proportions are different in each species, genus or group, and in particular in *S. ragei* n. gen., n. sp.: this derived “generalized anterior lobe scute pattern” is the intergular, which is wide and long, fully separates the short gulars and humerals, and widely reaches the pectorals. The particularities are here: 1) the position of the humeropectoral sulcus on the hyoplastron; 2) this close but posterior to the epi-hyoplastral suture; and 3) except reaching the lateral extremity of the suture on the left side. It is transversally positioned, following medially (intergular-pectoral sulcus part) to cross the entoplastron in the same direction, slightly posteriorly to the mid-length of

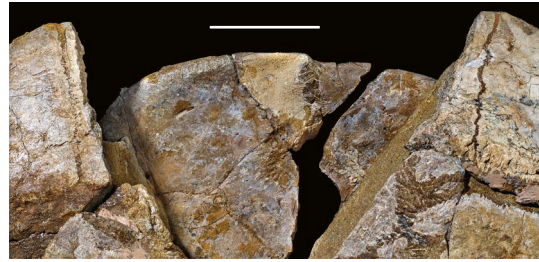


Fig. 7. — *Sindhochelys ragei* n. gen., n. sp., holotype CPAG-RANKT-V-3, from the Paleocene of Ranikot, locality K18-12, Southern Pakistan. Medioanterior part of the shell, the nuchal being removed, showing a part of the dorsal face of the anterior lobe with the sulci limiting the overlapping scales. Scale bar: 5 cm.

the plate; its direction is transverse as a whole and it is not incurved in a V medially on the entoplastron, contrarily to some Taphrosphyini including *Taphrosphys sulcatus* (in Hay, 1908), the *Carteremys* group and “*Carteremys*” *pisdurensis* Jain, 1977 (in Jain 1986) (Lapparent de Broin & Guntupalli 2020), and *Elochelys* (see Nopcsa 1931b; Gaffney *et al.* 2006); it is not laterally posteriorly directed towards the hyoplastral border contrarily to “*Carteremys*” *pisdurensis* in Jain (1986). *Sindhochelys ragei* n. gen., n. sp. characteristic gular-intergular pattern is: 1) the intergular is notably large; its precise borders are not well visible but its posterior width is estimated at its medial spring from the pectorals on the anterior two-thirds of the entoplastron; it is wider than each gular; it appears as wide all along the entoplastron, with possible parallel or nearly parallel lateral borders, but it becomes slightly wider between each gular on the epiplastra; and 2) each gular is shorter, and nearly reaches or not (at least on left side) the epientoplastral suture. The border extremities of the scutes is visible. This is seen (Figs 3; 6) at the incurved end of the intergular, gulars and humerals with their rounded border and by their ventral surface, which is slightly, individually, ventrally protruding (Fig. 6). The short plastral anterior scute overlap is visible on the dorsal face of the anterior lobe (Fig. 7). The posterior scute sulci are mostly visible. They are completed by symmetry on the Figure 4B. The pectoroabdominal sulcus is a little convex, positioned just anterior to the mesoplastra and reaching it, as seen on the left side. The abdominofemoral sulcus is also convex and its lateral left end indicates the bottom of the inguinal notch. The femoroanal sulcus is positioned on the xiphoplastron nearly parallel to the hypo-xiphoplastral suture, and it seems to be at mid distance from this suture and from the possible xiphoplastral extremity, if not slightly closer to the suture than to the anal notch.

The plastral scute length-formula is (medially except mediolaterally for humerals): femorals > pectorals > anals > abdominals = intergular > humerals > gulars.

Shell ornamentation (Figs 8; 9)

The whole shell surface is relatively damaged. As the turtle is very large, the general view does not show the surface details, so that we enlarged several small decorated parts representing the variability (Figs 8; 9AD). The shell was decorated by a net of meeting sulci delimiting irregular polygons, variable

in shape and proportions according to the position on the plates but relatively large on the whole, and protruded, i.e. much raised on the surface of the carapace and in relation to the sulci between the polygons. They are often damaged, their external surface being eroded, so that the complete depth of their protrusion (above the general surface) is not everywhere preserved (Fig. 8B, D). Some of the figures show imprints of the external surface in the matrix (Figs 8A; 9D). The polygons are either as wide as long or longer than wide. There are also places of short dichotomic sulci which are not united in polygons, and represent a usual basic decoration in aquatic turtles, but the elements have here a relatively marked length. In the right hyoplastron (Fig. 8C), there are short sulci, not united in a net, which follow the lines of the growing radiation of the pectoral scute (from the radiation center, situated towards the axillary buttress) (Fig. 3B). On the whole, on the external surfaces, the raised polygons are not regular and well delimited, and not as much protruding as in *Taphrosphys sulcatus* (Leidy, 1856), the type species of the genus (Fig. 9E-G). However, the less eroded part of the inner border of the peripherals 2 and 3 show smaller and well-delimited polygons 1 (Fig. 9B), and there are also such polygons on the suprapygal (Fig. 9C), which are rather more similar to those of *T. sulcatus*. But they do not constitute an as well constituted net of granulations protruding on the whole shell surface.

COMPARISONS

General comparisons of the carapace of Sindhochelys ragei n. gen., n. sp. with Bothremydidae

The carapace of *Sindhochelys ragei n. gen., n. sp.* differs from that of other bothremydids in a combination of unique features: 1) it has no parallel lateral borders in its middle part; 2) it is rounded as in various other bothremydids but differently by the position of its largest width at peripherals 6-7 sutures and not posteriorly; 3) together with an anterior rounded border anterior to this line; border that was straight just at nuchal border and probably the medial part of the peripheral 1, but not making a medially protruding carapace at nuchal area; and 4) and together with the slightly pointed in V posterior border. All that makes the carapace neither rectangular nor ovoid. It is distinct from the more rectangular to ovoid shape with long parallel median lateral borders of various other bothremydid forms.

However, some features of *S. ragei n. gen., n. sp.* are distributed in mosaic in various bothremydid species. The shape of the carapace of *S. ragei n. gen., n. sp.* shows similarities with the rounded shell of the holotype of the bothremydine *Chedighaii barberi* (Schmidt, 1940) (Gaffney & Zangerl 1968) from the late Cretaceous (Campanian) of Arkansas (United States); in this form, the lateral profile shows a moderate height and much oblique bridges (Schmidt 1940: figs 3 and 4) as, for example, in *Francemys* (from Gadoufaoua) and *Eusarkia rotundiformis* Bergounioux, 1952 – “*Gafsachelys*” group (a Tunisian Ypresian taphrosphyine); such relatively high inclined bridges were probably present in *S. ragei n. gen., n. sp.* because of the wide bridge processes and the large width

of the shell, making it rather low (Fig. 2); but its inclination angle also exists in other wide bothremydids shells. The bothremydine species *Ch. barberi* is also figured under the name *Podocnemis alabamae* Zangerl, 1948 (Zangerl 1948: fig. 4.1) (Campanian) (a junior synonym, see Gaffney *et al.* 2006), by a specimen which differs in a slight intraspecific variability in width, usual in turtles: some individuals are slightly wider than others, thus appearing shorter, which is an intraspecific difference known as sexual dimorphism in living forms. Both *Ch. barberi* specimens are large individuals (maximal lengths between 58.2 and 65 cm). But in this species, the widest margin is at the peripheral 7 (posterior to the suture 6-7) and the anterior peripherals and nuchal are less elongated and wide. Beside this difference, *S. ragei n. gen., n. sp.* differs in the less medially inclined borders of peripherals 2 and 3, an anterolaterally wider costal 1 and a longer costal 1 + neural 1 + nuchal part; probably it differs also in the absence of a nuchal notch, on a nuchal that is narrower for its length and anteriorly relatively wider in *S. ragei n. gen., n. sp.* *Sindhochelys ragei n. gen., n. sp.* has lateral borders of plastral lobes less oblique; the anterior lobe is relatively slightly shorter and anteriorly wider, being more trapezoidal with lateral borders less converging anteriorly, and the posterior lobe is less wide: all that is seen when the carapaces are adjusted in width and length and superposed. The shell of *Ch. barberi* (from these two specimens) is more similar to that of another bothremydine, *Rosasia soutoi* Carrington Da Costa, 1940, from the late Cretaceous (Campanian) of Portugal (Antunes & Broin 1988), sharing a shorter carapace anterior part than in *S. ragei n. gen., n. sp.*, and an anterior border with a notched and anteriorly narrower nuchal; the carapace of *R. soutoi* is more regularly rounded than in *Ch. barberi*, with a short median length of parallel lateral borders.

By its large size and Paleocene age, the carapace of *Sindhochelys ragei n. gen., n. sp.* can be compared with the large carapace of *Puentemys mushaisaensis* Cadena, Ksepka, Jaramillo & Bloch, 2012, from the Paleocene Cerrejón Formation (Colombia) (maximal carapace length: 118 cm). But the latter has differently inclined borders of peripherals 2 and 3, because of an overall medially elongated and protruding nuchal part, giving a more elongate and more oval anterior carapace. This shape differs from what is observed in *R. soutoi* and *Ch. barberi* (Bothremydini) and *S. ragei n. gen., n. sp.*, as well as from that of *Foxemydina* Gaffney *et al.* 2006; this is a late Cretaceous European infratribe in which *Puentemys* has been included by Cadena *et al.* (2012). *Rosasia soutoi* and *P. mushaisaensis* have 7 neurals as in *S. ragei n. gen., n. sp.*. But in *P. mushaisaensis*, these neurals are much wider, as such is also the vertebral 1 anteriorly, which reaches the peripherals 2. Besides, the plastral lobes of *P. mushaisaensis* are particularly distinctive. Its anterior lobe is relatively longer and anteriorly narrower (in agreement with the dorsal carapace border), whereas the posterior lobe is wider and much more rounded, filling the opening space of the posterior carapace including the inguinal notches.

The bothremydines *R. soutoi*, *Ch. barberi* and *P. mushaisaensis* surely do not belong to the group of *Sindhochelys* because,

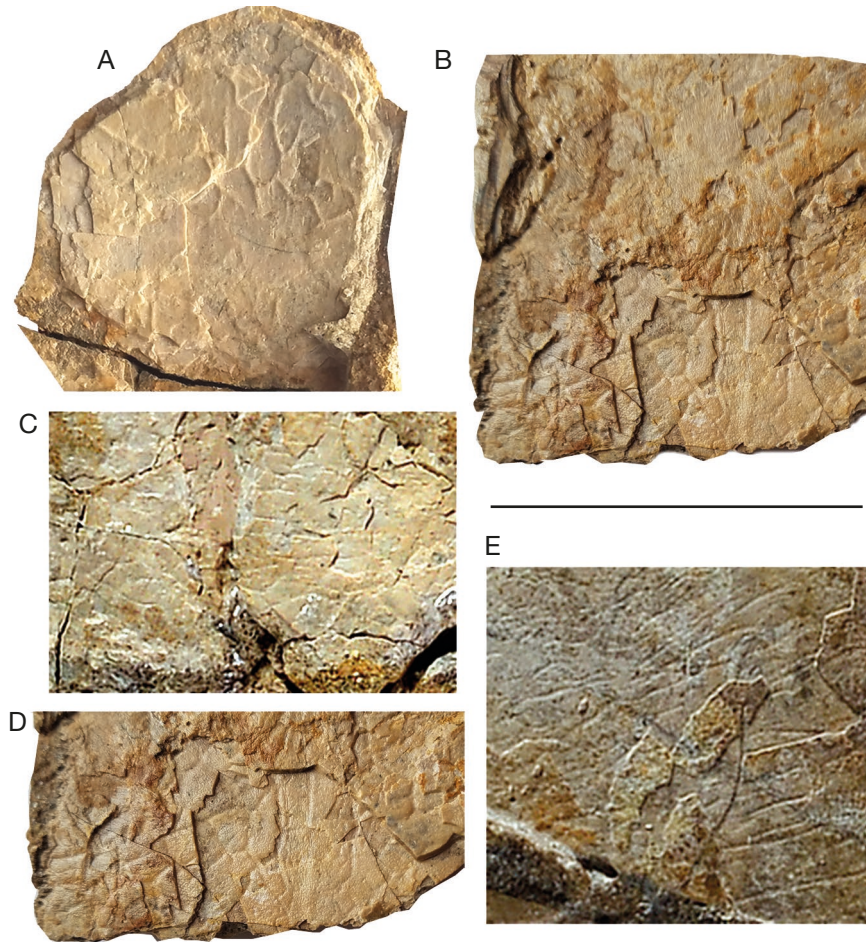


FIG. 8. — *Sindhochelys ragei* n. gen., n. sp., holotype CPAG-RANKT-V-3, from the Paleocene of Ranikot, locality K18-12, Southern Pakistan. **A-D**, decoration in various parts of the shell: **A**, isolated piece with the imprint of the mesoplastral external surface; **B**, marginal part of peripheral 10, dorsal view; **C**, left costal 1 part, posterior border at mid-width, dorsal view; **D**, a costal lateral border, in dorsal view; **E**, right hyoplastral part, ventral view. Scale bar: 5 cm.

beside their own derived particularities, they differ from *S. ragei* n. gen., n. sp. in primitive features, which are also present in many other bothremydids (features which are derived in *S. ragei* n. gen., n. sp.): 1) the primitive plastral anterior scute pattern with an intergular separating the smaller gulars, longer than them but not reaching the pectorals, so that the humerals meet medially; 2) the absence of any well marked polygonal ornamentation. *Chedighaii barberi* ornamentation is described thus (Gaffney *et al.* 2006): “Shell surface texture is the “pelomedusoid” pattern (Broin, 1977)” (p. 551) “consisting of fine forking and irregular vascular grooves” (p. 548), i.e. the basic decoration seen in aquatic forms. *Rosasia soutoi* also presents a basic decoration (Carrington da Costa 1940: fig. 2). In *P. mushaisaensis*: “The carapace is smooth, lacking decoration on the surface in the holotype and all referred specimens” (Cadena *et al.* 2012: 691). All the other species presenting together the basic decoration (more or less marked) – or lacking any description and figure of the decoration – and a primitive plastral scute pattern, have a more different contour of carapace and plastron, even if many other details may be similar and distributed in mosaic in others, such as the elongation degree of the anterior cara-

pace border, for example. Some specimens of a species have a more marked decoration than others, such as the specimen from the Cenomanian of Portugal attributed to *Algorachelus peregrinus* Pérez-García, 2016 by Pérez-García *et al.* (2017a): such a decoration is not visible in the figured type series specimens of *A. peregrinus* Pérez-García, 2016 from Spain (Pérez-García 2018a). *Elochelys perfecta* Nopcsa, 1931 (France) (two known specimens) has also, similarly to this Portuguese *Algorachelus* specimen, a more marked basic decoration of mostly elongated polygons, which are not raised in relation to *S. ragei* n. gen., n. sp. The nuchal of an undetermined Pelomedusoides from the Eocene of the Gujarat Province (India) (Smith *et al.* 2016) presents such a well-marked basic decoration of not raised small polygons. However, unlike in *Algorachelus*, *Chedighaii*, *Rosasia* and *Puentemys*, *E. perfecta* shares with *Sindhochelys* the derived “generalized plastral scute pattern”, defined by an elongated intergular, fully separating the humerals up to the pectorals. However, within this frame, *E. perfecta* differs from *S. ragei* n. gen., n. sp. in the anterior position of the humero-pectoral sulcus. In *E. perfecta*, it is anterior to the epihyoplastral suture and, medially the intergular is V shaped on the entoplastron, with the V

point posterior to the epihyo-plastral suture level: that makes a clearly distinct contour and proportion of the intergular in both species. Besides a moderate decoration and the different intergular shape, the shell shape of *E. perfecta* differs from that of *S. ragei* n. gen., n. sp. in a suite of features such as the anteriorly and posteriorly more rounded carapace, with parallel lateral borders on a small median length and with an anterior rounded protrusion, and a different plastron by the more rounded anterior plastral lobe, and the wider and more rounded posterior lobe.

Particular comparisons of Sindhochelys ragei n. gen., n. sp. with some Indian Bothremydidae

Alone in India, two Indian bothremydid species share one or two of the most important general derived features of *S. ragei* n. gen., n. sp.: the generalized anterior lobe scute pattern as in *E. perfecta* (in the above compared forms) and the protruding polygonal ornamentation (unlike the above compared forms).

1) *Carteremys leithii* (Carter, 1852), sharing both features with *Sindhochelys ragei* n. gen., n. sp. *C. leithii* from the Upper Cretaceous (Maastrichtian) intertrappean-beds of the Worli Hills (Mumbai, India) has been revised and figured by a specimen (GSI coll. no. 20337) in Williams (1953: pl. 3). This has been designed as neotype in Lapparent de Broin & Guntupalli (2020). Williams's photograph is not fully informative but the decoration is visible on the figure and described by Williams (1953: 6) as follows: "the characteristic sculpture of the surface (mentioned by Carter), which, while somewhat like that of some chelyid, also resembles that of, for example, the American pelomedusine genus *Taphrosphys*". This sculpture particularly appears on his figure, on the left costal 1, as made of small polygons that are well protruded and regularly elongated: the polygons are more regular and smaller than in *S. ragei* n. gen., n. sp. They appear as more similar in outline to those of *Elochelys perfecta* but differ in the well-marked protrusion. The plastron of *C. leithii* is known by an incomplete drawing of Carter (1852), missing the sutures of the anterior lobe plates. The anterior plastral scute pattern has similarities with that of *S. ragei* n. gen., n. sp. This is particularly visible in the long intergular separating the humerals and reaching the pectorals. However, the intergular contact with the pectorals is just marked by a point in the drawing. This short punctiform contact is rather unusual; it is known to occur as a variable feature in some specimens of the podocnemidid *Neochelys arenarum* Broin, 1977, beside a wider contact in other specimens. However, other specimens of the *Carteremys* group from India complete the shape variability for the species group, with the intergular having a wider contact with the pectorals (Lapparent de Broin *et al.* 2009; Lapparent de Broin & Guntupalli 2020). The shape of the carapace of *C. leithii* (as revised by Williams 1953) is very distinctive from that of *S. ragei* n. gen., n. sp. and *E. perfecta*: it is anteriorly narrower, with a different proportion of enlarged peripheral part and nuchal, and with the widest carapace part at the 7th-8th

suture, which is posterior to the peripherals 6-7 contact of *S. ragei* n. gen., n. sp.

and 2) "*Carteremys*" *pisidurensis* Jain, 1977, sharing the generalized scute pattern with *Sindhochelys ragei* n. gen., n. sp. "*C.*" *pisidurensis* from the Upper Cretaceous (Maastrichtian) Lameta Formation (infratrappian beds) of (India) also has a similarly generalized scute pattern on a drawing of Jain (1986: fig. 3, for "*Shweboemys*" *pisidurensis*). But the humero-pectoral sulcus is just anterior to the epihyo-plastral suture level, medially on the entoplastron, and it becomes posterior and oblique towards the external hyo-plastral border, while the intergular shape is posteriorly wide in a V on the entoplastron: the drawing of the sulci is not visible on the photographs, and the drawing of the carapace is problematic on the nuchal part (the form needs to be reexamined to rename the genus, see Lapparent de Broin & Guntupalli 2020). Contrarily to *Sindhochelys ragei* n. gen., n. sp., the shell is not decorated and the shape of the carapace is clearly not similar: on the photographs, the carapace is quadrangular and not oval, medially widened toward the midline, contrary to Jain's drawing.

Particular comparisons of Sindhochelys ragei n. gen., n. sp. with Taphrosphyini

The members of the tribe Taphrosphyini Gaffney, Tong & Meylan, 2006, emend. Lapparent de Broin & Guntupalli (2020) present the two characters shared by the *Carteremys* group beside differences which characterize the Taphrosphyini tribe. In this work, this tribe is restricted to the *Taphrosphys* group of Broin (1988) in Antunes & Broin (1988), contrarily to Gaffney *et al.* (2006) and Pérez-García (2019b). All the species included by Gaffney *et al.* (2006) in their "taphrosphyine" tribe which have no *Taphrosphys* decoration, carapace shape other than that of *Taphrosphys*, and presenting other different characteristics of skull, are united in the tribe "Nigeremydini Gaffney *et al.*, 2006" new rank in Lapparent de Broin & Guntupalli (2020) (i.e. basically the *Nigeremys* group of Broin in Antunes & Broin [1988]) (Lapparent de Broin *et al.* 2020). Besides their different anatomy, by their stratigraphical first appearance as soon as late Cretaceous and wide geographical diversification, the Taphrosphyini of Gaffney *et al.* 2006 (emended) cannot be the sister group of the *Azabbaremys-Acleistochelys* clade of Gaffney *et al.* 2006 (Gaffney *et al.* 2001c, 2006, 2007), a Malian Paleocene clade which is the youngest element of Nigeremydini. This tribe is a group of large turtles living in the Maastrichtian-Paleocene Trans-Saharan seaway, extending to Neotethys as a gulf or a full passage (according to the age) in the African craton (Halstead 1979; Walker 1979; Lapparent de Broin & Werner 1998; Lapparent de Broin 2000; Lapparent de Broin & Guntupalli 2020; Lapparent de Broin *et al.* 2020). Thus, newly understood and emended, Taphrosphyini tribe is represented by the type species *Taphrosphys sulcatus* (Leidy, 1856) from the early Paleocene (Danian) of New Jersey, although the tribe is already widely diversified at Maastrichtian times (Africa, Syria, France). It is defined by shell features such as: 1) the autapomorphic long pubic scar and small rounded

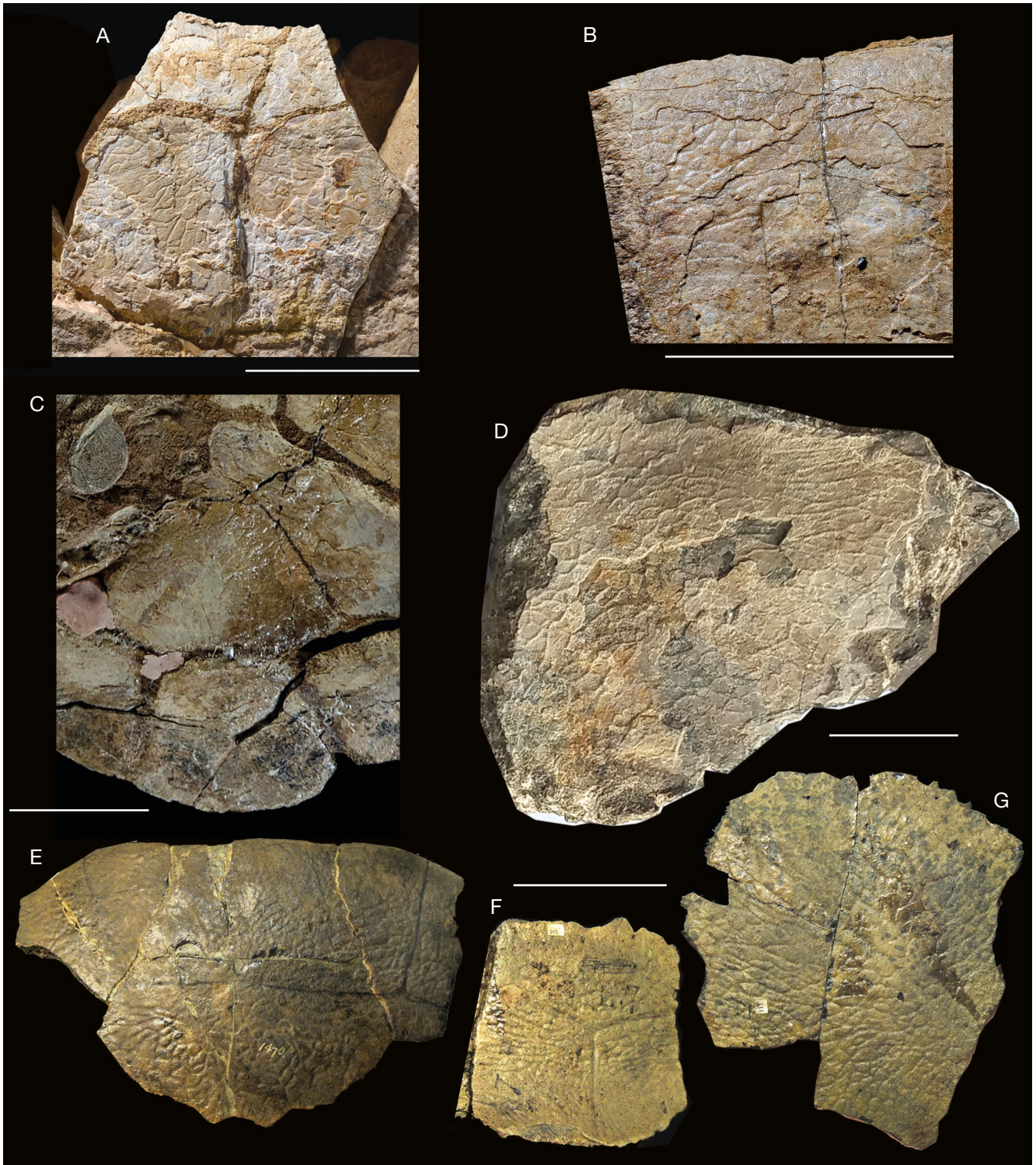


FIG. 9. — Comparison of the decoration of *Sindhochelys ragei* n. gen., n. sp. from the Paleocene of Ranikot, locality K18-12, Southern Pakistan, and *Taphrosphys sulcatus* (Leidy, 1856) from the early Paleocene of New Jersey (United States): **A-D**, *Sindhochelys ragei* n. gen., n. sp., holotype CPAG-RANKT-V-3: **A**, nuchal part; **B**, left peripherals 2, dorsal face with smaller polygons than in other parts; **C**, suprapygals-pygals part, dorsal views; **D**, imprint of an isolated plate, ventral view; **E-G**, *Taphrosphys sulcatus*: **E**, AMNH 1470, “Barnsboro, N.J.” (Cope collections), nuchal border of three plates, dorsal view; **F**, **G**, AMNH 2522, “Tinton Falls, N.J.”, syntype: **F**, peripheral 6 (medial plate of three in Leidy (1856: pl. 19, fig.4), and in Gaffney *et al.* (2006: fig. 20), dorsal view; **G**, xiphiplastron, ventral view. Scale bars: A-C, E-G, 5 cm; D, 2 cm.

or triangular spot of the ischiatic scar (which is situated at the border of the anal notch of the xiphiplastron) (Gaffney 1975: fig. 11; Hay 1908: fig. 118; Pérez-García *et al.* 2018);

and 2) to which is obligatorily added the regular decoration of protruding small polygons (Fig. 9E-G) and the generalized scute pattern of the anterior lobe.

Taphrosphys sulcatus (Leidy, 1856) representative of the *Taphrosphyini* tribe

Thus, all the referred specimens to *Taphrosphys* have the generalized anterior lobe scute pattern (above defined), like that of *Sindhochelys ragei* n. gen., n. sp. The syntype of *T. sulcatus*, AMNH 2522 (under the name of *Platemys sulcatus* Leidy, 1856) is of unknown locality and horizon (“Tinton Falls, N. J.” in Leidy 1865: 109), without any further information except that it belongs to the Hornerstown Formation, which was first attributed by Gaffney (1975) to the late Cretaceous. The syntype consists of a fragment of 5th, 6th, and 7th left peripherals (Leidy 1865: pl. 19, fig. 4, a drawing reproduced in Gaffney *et al.* [2006], fig. 20, reproduced in part in Fig. 9F) and a left xiphiplastron (Fig. 9G). Gaffney *et al.* (2006) noticed that if only the peripherals were figured by Leidy, it is the xiphiplastron that has the diagnostic features of the genus and it may become the lectotype. Moreover, the fossil MFL area of the Hornerstown Formation that yielded all the *T. sulcatus* specimens is now considered as basal Danian (Gaffney *et al.* 2006), as is *S. ragei* n. gen., n. sp. No more photographs of the shell were given except one photograph of a dorsal incomplete shell (specimen PU 18707, from Sewell [N. J.] in Gaffney [1975: fig. 5]) with a reconstructed contour; the polygonal ornamentation, similar to that of the Fig. 9E-G, is barely visible, as well as on Gaffney’s figure 5 as on the field photograph of the dislocated specimen PU 18707 in situ. For the reconstruction of the carapace shape of *T. sulcatus*, beside that of PU 18707 (Gaffney 1975), we consider the figures of Hay (1908) from various incomplete New Jersey specimens, which were attributed to various species including the species *T. leslianus* Cope, 1870, *T. (Prochonias) longinuchus* (Cope, 1870) and *T. molops* Cope (1870) (all species synonymized with *sulcatus* in Gaffney [1975]). Among others, Gaffney *et al.* (2006) mentioned the specimen AMNH 1470, from Barnesboro (N. J.), which notably includes the anterior carapace part, described as *T. molops* by Hay (1908: 119) and figured here (Fig. 9E). As the syntype, it shows the specific ornamentation, and besides, a contour of the anterior margin of the shell that conforms that of the other preserved specimens, as figured in Hay (1908) and preserved in the AMNH fragments.

Comparison of the ornamentation of Sindhochelys ragei n. gen., n. sp. with *Taphrosphyini*

The comparison of the nuchal ornamentation of *T. sulcatus* with that of *Sindhochelys ragei* n. gen., n. sp. shows a rounded granulation of smaller and more regular polygons (Fig. 9E-G compared with Fig. 9A-D and Fig. 8). Open dichotomic sulci of *S. ragei* n. gen., n. sp. carapace parts (Fig. 8C, E) are not known in the *T. sulcatus* assemblage of photographed specimens. Various elements of carapace and plastron of other *Taphrosphyini* present a decoration similar to that of *T. sulcatus* and are known by photographs, such as *T. congolensis* Wood, 1975 from the Paleocene of Angola (Wood 1975; Pérez-García *et al.* 2018) (our observation at MRAC). Other *taphrosphyine* forms may also include raised polygons and dichotomic sulci at some places, such as *Eotaphrosphys ambiguus* (Gaudry, 1890) from the Maastrichtian of Mont-Aimé (France)

(MNHN.F.MTA1) (Broin 1977; Montenat & Merle 2018: fig. 164B1, D1; Pérez-García 2018b), which displays some similarities with those of *S. ragei* n. gen., n. sp. (Figs 3A; 8; 9A-D). Well raised irregular and regular polygons are present in another *taphrosphyine*, the *Eusarkia*-“*Gafsachelys*” group of the Ypresian of Gafsa (Tunisia) (Bergounioux 1952, 1956). There also, on the whole a more evident net is always present. The polygons radially diverge from a scute center, to become elongated on the borders, as in *S. ragei* n. gen., n. sp., *T. sulcatus* elements (Fig. 9E-G) and *Eotaphrosphys ambiguus* for example. This radiation is visible in *S. ragei* n. gen., n. sp. on the nuchal and suprapygal (Fig. 9C), but the polygons differ in size and shape. This Tunisian group is an assemblage of undefined number of “species” (see a discussion in Moody (1972), Broin (1977), Gaffney *et al.* (2006)). It was established on several individuals where the decoration has its variabilities, beside the variable medial union of the costals over the neurals but with the constant medioanteriorly protrusion of the ovoid carapace, the longest known protrusion in the tribe. The group includes *Eusarkia rotundiformis* Bergounioux, 1952 (Bergounioux 1952: pl. 1-2) (holotype MNHN.F. coll.) mostly smoothened surface by erosion and lacking the anterior border, *Gafsachelys neurriregularis* Bergounioux, 1952 (pls 3-4) and Bergounioux 1956 (pls 10[2], 11, 12 and 13), *G. moulaensis* Bergounioux, 1956 (Bergounioux 1956: pl. 10[1]), and *Euclastochelys interrupta* Bergounioux, 1956 (Bergounioux 1956: pls 14, 15), all variably preserved. The part of shell named *Gafsachelys phosphatica* Bergounioux, 1952 (ENSMP, MNHN.F deposit) is a *nomen dubium*. It cannot be the type specimen of the genus (generotype) and the type species of the assemblage. So that, one other species ought to be retained for the whole assemblage (see Broin 1977) and this might be the holotype shell of *Eusarkia rotundiformis* Bergounioux, 1952 although it does not present all the diagnostic characters of the species. In these Tunisian specimens, the polygons are protruding, when they are not mostly eroded as in *Eusarkia rotundiformis*, and they form, as in *Taphrosphys sulcatus*, a better defined net of polygons than in *S. ragei* n. gen., n. sp. During the Maastrichtian of other countries, some isolated fragmentary specimens individually preserve diagnostic features of *T. sulcatus*: 1) xiphiplastral shape and scars; 2) similar anterior lobe scute sulci; 3) similar anterior elongation and rounded anterior shape; 4) characteristic ventral scars on the costal 1 of the hyoplastral buttress and ribs 1 and 2; and 5) short costals, posterior to the first one. They have a similar fine and regular protruding ornamentation as in *T. sulcatus*; the small polygons (which are elongated or not according to their place on the plates) are well united in the same net as in *T. sulcatus*, *T. congolensis* and the Tunisian group, contrarily to the *S. ragei* n. gen., n. sp. ornamentation. These Maastrichtian remains are attributed to *T. sulcatus* in phosphates of Syria, at Charquieh (Bardet *et al.* 2000: figs 7a, b, c) and to *Taphrosphys* cf. *sulcatus* in Egypt, at Ammonite Hills, close to the Dakhla oasis (Lapparent de Broin & Werner 1998: fig. 3 [3]). Two species of *Taphrosphys* from the Maastrichtian of Mont-Aimé (France) and Eocene of Peru (respectively) have been separated from the genus and placed in two new genera

by Pérez-García (2018b) and Pérez-García *et al.* (2018): they share *Taphrosphys* principal diagnostic features (decoration, plastral scute pattern and various others). The Maastrichtian Syrian and Egyptian shell elements which are the closest to *T. sulcatus* among the Taphrosphyini have not been integrated in these works, no more than new Maastrichtian taphrosphyine carapaces of Phosphates of Morocco (Bardet *et al.* 2018) (OCP collections observed at Khouribga): this Maastrichtian presence shows that the tribe Taphrosphyini is recorded by *Taphrosphys* or cf. *Taphrosphys* in Africa and Syria and by a close taxon in France as early as the Maastrichtian. Thus, during the Maastrichtian, the tribe occupied a widespread distribution along the northern platform coast, Neotethyan borders and appendices, including its presence as northern as at the Mont-Aimé (France) and western in Morocco. The tribe is well defined as a derived monophyletic group before its Paleocene radiation, before the derivation of the nigeremydine *Azzabaremys-Acleistocheilus* clade and before the presence of *S. ragei* n. gen., n. sp. in Pakistan, which does not appear to be associated with the taphrosphyine tribe.

Comparison of the shape of the carapace of Sindhochelys ragei n. gen., n. sp. with Taphrosphyini

Although the ornamentation shows some similarities, the comparison of the shell shape of *Taphrosphys sulcatus* with that of *Sindhochelys ragei* n. gen., n. sp. does not support a close relationship between them. The carapace may not be as important for its length as that of *S. ragei* n. gen., n. sp., even taking into account a possible relative difference of width or length due to the sexual dimorphism: the shape variability in *T. sulcatus* is shown by the figures of Hay (1908) as limited in one specific frame. The carapace of *T. sulcatus* is reconstructed in Hay (1908) and Gaffney (1975) by different contours and some of the Gaffney's figures need a revision. As seen in Figure 9E and in figures of Hay (1908), even in the more anteriorly protruded carapace, the anterior border is not the most protruded at the nuchal alone as figured in two reconstructions of Gaffney (1975: figs 1 and 5): these two figures give the species a much ovoid shape; but the carapace anterior border is straight at the medial part of the peripherals 1 beside at the nuchal border. It is the same even in the more taphrosphyine protruded carapaces, those of the species of the *Eusarkia*-“*Gafsa*” group (see “*Gafsa*” *neurriregularis*” Bergounioux, 1952 and “*G. moularensis*” Bergounioux, 1956). In *T. sulcatus*, laterally to this anterior straight border, the borders diverge posteriorly as figured in Gaffney (1975: fig. 5). From all the partial specimens figured in Hay (1908), the anterior curve of the contour is too much rounded in the two reconstitutions of Gaffney (1975: figs 1, 5). As a whole, the carapace of *T. sulcatus* is overall ovoid, although not completely rounded at the anterior medial part, with the posterior widening at the peripherals 7-8 (presenting lateral borders parallel on that short length), followed by a progressive posterior curved protrusion, towards a median protrusion that is maximal at the level of the pygal, as presented in Gaffney (1975: fig. 5); but the carapace does not present a posterior trapeze with rounded angle as in Gaffney (1975:

figs 3 and 4): his best reconstruction ought to be his figure 5, adding a correction for a narrow straight border at the nuchal and medial part of the adjacent peripherals 1. Endly, the best precise carapace representation of *T. sulcatus* appears in the figure of the type of “*Taphrosphys longinuchus*” Cope, 1870, in Hay (1908: fig. 101). As anteriorly described in this way, with its posteriorly positioned maximal width and with the not in V pointed posterior border, the carapace is clearly different from that of *S. ragei* n. gen., n. sp.. The shape of the carapace of *T. congolensis*, as reconstructed by Pérez-García *et al.* (2018), appears as morphologically close to that of *T. sulcatus*, differing by the small nuchal notch of the former, and knowing the plastra are globally similar (Gaffney 1975; Wood 1975; Pérez-García *et al.* (2018).

The differences between *Taphrosphys* and *Sindhochelys ragei* n. gen., n. sp. are also those that differentiate the latter from *Carteremys leithii*. In *C. leithii*, the anterior part of the carapace is not as elongate as in *T. sulcatus* and not as wide as *S. ragei* n. gen., n. sp., and *C. leithii* does not appear as being a taphrosphyine (see Lapparent de Broin & Guntupalli 2020). In other Taphrosphyini species, the carapace is not enough complete but when preserved, the shape is likewise mostly ovoid as that of *T. sulcatus*. Among the Tunisian specimens from Gafsa, the best specimens, with a nearly complete shape, are “*Gafsa*” *moularensis*” and the similar “*G. neurriregularis*” in Bergounioux (1956: pl. 10). In these individuals, the anterior protrusion is preserved: it is specifically maximal in relation to other taphrosphyines, with a narrower nuchal-peripheral 1 part – anterior straight border and they have less medially inclined and longer lateral borders, framing a very elongated medial part; so that the carapace looks more strictly “ovoid,” being more protruded than in *T. sulcatus*, and anyway far from the contour of *S. ragei* n. gen., n. sp. However, Gafsa form shape remains in the general frame of that of *T. sulcatus*.

Comparison of the plastral shape of Sindhochelys ragei n. gen., n. sp. with Taphrosphyini

In addition to the decoration, the comparison of the scute pattern of the anterior plastral lobe shows that all members of Taphrosphyini have the generalized anterior lobe scute pattern reconstructed for *Sindhochelys* (Figs 3B; 4B), i.e. that of a wide and long intergular separating the humerals and contacting the pectorals, and, further, the humeropectoral sulcus is not anterior to the epihyoplastral suture in Taphrosphyini and *S. ragei* n. gen., n. sp. The complete plastron of *T. sulcatus* is not fully preserved on each specimen, and the reconstruction (Gaffney 1975) did not include the variabilities presented by other New Jersey specimens in Hay (1908): the position of the humeropectoral varies from linked to the epihyoplastral suture in “*T. molops*” (type AMNH 1472), slightly more than in *S. ragei* n. gen., n. sp., to far posterior in “*T. leslianus*” (AMNH 1471, a specimen with the epiplastron and a left hyoplastron but without the entoplastron). Thus, this sulcus is never anterior to the epihyoplastral suture as in other taphrosphyines (Montenat & Merle 2018: fig. 164B1; Pérez-García *et al.* 2018). The entoplastron is figured by Hay (1908) as variably long posterior to this suture. At the differ-

ence from *S. ragei* n. gen., n. sp., the intergular is pointed in a medial V, at the junction with the pectorals (as in *Elochelys perfecta*) in “*T. molops*” figured specimen of Hay (1908); but the V is variable in length in other “species”, where the sulcus is, nevertheless, not as transversal as in *S. ragei* n. gen., n. sp. and in some *Carteremys* group members (Lapparent de Broin *et al.* 2009; Lapparent de Broin & Guntupalli 2020). The scute plastral pattern of *T. sulcatus* is the same in other Taphrosphyini, including the *Eusarkia*-“*Gafsachelys*” group: the sulci are not correctly figured or visible in all drawings and photographs of Bergounioux (1952; 1956), but the V is visible on “*Gafsachelys neurriregularis*” (Bergounioux 1956: pl. 123, fig. 1), and the holotype of *Eusarkia rotundiformis*, with the V pointed intergular.

The plastron of *Sindhochelys ragei* n. gen., n. sp. has not only a similar generalized scute pattern (although with the above exposed differences concerning the sulcus between humerals, intergular and pectorals precise position), but it has another similarity with *T. sulcatus* by the trapeze shape of the anterior lobe, although it is wider in *S. ragei* n. gen., n. sp. In fact, as *S. ragei* n. gen., n. sp. has its own proportions, beside the “species of Hay, 1908” assembled in *T. sulcatus*, each other taphrosphyine species has its own proportions in the width of the lobes, shape of the lobes, degree of convergence of the lateral borders, proximity of the abdominofemoral sulcus with the hyohypoplastral suture, width of the posterior lobe at abdominofemoral sulcus in relation to the anterior lobe at the bottom of the axillary notches, and degree of rounding of the posterior lobe posteriorly to this sulcus: because of individual small variations, with only one specimen by species, it is difficult to define the characteristic proportions of each species in a defined specific frame. For the posterior lobe shape, two taphrosphyines (each one preserved by one specimen) share with *S. ragei* n. gen., n. sp. (one specimen) a posterior lobe with few or barely rounded lateral borders: *T. congolensis* and *Eotaphrosphys ambiguus* (Gaudry, 1890) (Broin 1977; Montecat & Merle 2018; Pérez-García 2018b; Pérez-García *et al.* 2018). *T. sulcatus*, as it is reconstructed in Gaffney (1975), on the base of the syntype xiphiplastron and on the other New Jersey fragmentary specimens (Hay 1908), has a posterior lobe anal part with a slightly concave lateral border and it is more or less narrowed: that has a possible sexual variability. *Motelomama olssini* (Schmidt, 1931) from the Eocene of Peru, and the Ypresian *Eusarkia*-“*Gafsachelys*” group from Gafsa (Bergounioux 1952, 1956), share a wider posterior lobe at its base and also a narrowed rounded anal part. As far as *S. ragei* n. gen., n. sp. is concerned, the posterior lobe has a weak progressive convexity of the lateral borders, which is similar to that of *Eotaphrosphys ambiguus*: this is a less derived feature than a wide and more rounded lobe border and a more narrowed anal part, as present in the other above cited species. In a cladistic analysis, it simply represents a primitive character and not a derived shared character by *S. ragei* n. gen., n. sp. and *T. sulcatus*. The taphrosphyine anal notch is differently deep, rounded or straight and wide or narrow, often concave with pointed

extremities, varying in each species: it varies from primitively wide and very short in *E. ambiguus* (Montecat & Merle 2018: fig. 164C), slightly more concave (and variably wide) in *T. sulcatus* and up to narrower and longer in *M. olssini*. The anal notch of *S. ragei* n. gen., n. sp. is incomplete: from the remaining border, it was probably morphologically close to that of *E. ambiguus* in length and width, possibly not concave with not pointed ends, and then probably relatively primitive among bothremydids. The inner face not being visible in *S. ragei* n. gen., n. sp., it cannot be appreciated if there was a taphrosphyine ischiatic scar with its characteristic rounded spot; the extremity of the plastron is broken at the ischiatic suture, and, looking in posterior view, it does not seem the spot was present.

Finally, although similarities with bothremydid species, which share the same “generalized anterior lobe scute pattern” with a weak (*Elochelys perfecta*) to more strongly protruded (*Carteremys leithii* group, Taphrosphyini) decoration or without a decoration (“*Carteremys*” *pisduensis*), *Sindhochelys* is different from all bothremydids together by the shape of the shell, its decoration and its unique scute/suture relations, particularly in the anterior lobe scute pattern.

Gen. et sp. indet.
(Fig. 10)

EXAMINED MATERIAL. — Fragment of a carapace constituted of two right peripherals with adjacent peripherals and costal parts, CPAG-RANKT-V-4.

LOCALITY. — Lower part of the Khadro Formation of the Ranikot Group, early Paleocene, locality K18-12, in the proximity of Ranikot Fort, Jamshoro District, Sindh Province, Southern Pakistan (Fig. 1).

DESCRIPTION AND COMPARISONS

The specimen (17 cm long × 10 cm wide) includes a right bridge portion of carapace with the posterior part of the peripheral 5, the complete peripherals 6 and 7, with a small part of the peripheral 8, sutured to the lateral end of the costals 3 (part), 4 (complete extremity) and 8 (minimal part). In dorsal view (Fig. 10B, C), the peripherals (complete 6th and 7th) are wide and display an incurved surface. The free border is thick but neither rounded nor acute. In ventral view (Fig. 10A), the posterior extremity of the bridge is positioned at first third of the 7th peripheral. The sutured border of the plates with the plastron is weakly indented to receive the corresponding weak indentations of the hypoplastron. The fragment was colonized by a perforating animal, such as the pholad bivalve, which nested forming rounded cavities. That occurred once the carapace was dislocated in the marine-littoral water ground, because the cavities are distributed as well on the external face, on the periphero-hypoplastral suture and in the inner bridge cavity, being of various sizes.

The fragment is noticeable by its ornamentation, which does not substantially differ from that of *S. ragei* n. gen., n. sp., being made of wide protruding polygons (Fig. 10C)

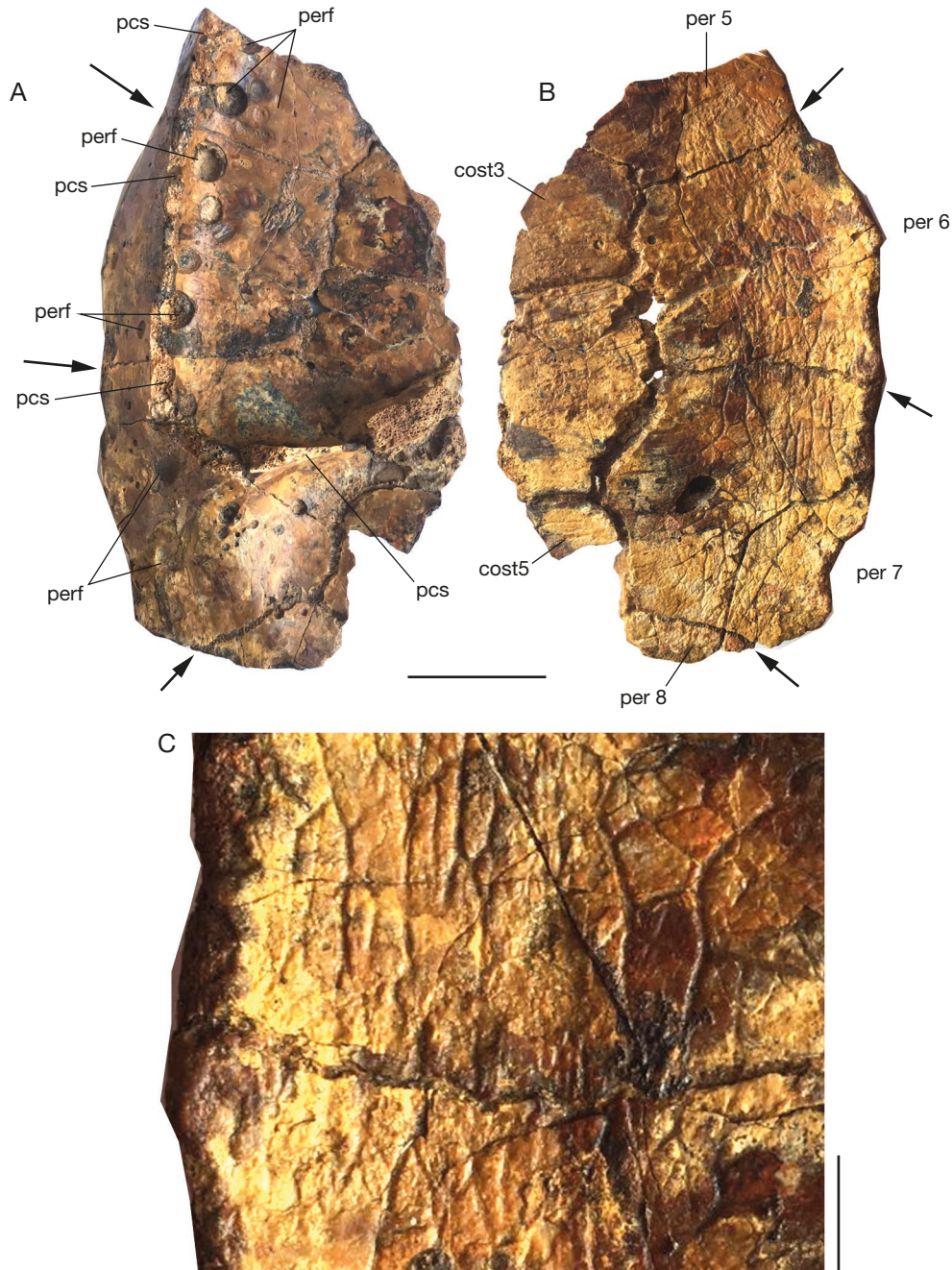


FIG. 10. — Fragmentary part of a carapace of an undetermined Bothremyidae, from the Paleocene of Ranikot, locality K18-12, CPAG-RANKT-V4, Southern Pakistan: **A, B**, ventral and dorsal views; **C**, detail of the dorsal decoration. Abbreviations: **cost3**, costal 3; **cost5**, costal 5; **pcs**, suture of the peripheral border with the border of plastral processes; **per 5-8**, peripherals 5, 6, 7, 8; **perf**, pholad perforations. Scale bars: A, B, 4 cm; C, 1 cm.

but which are more marked. The polygons are very elongated and well defined and as elongated as in some plates of *S. ragei* n. gen., n. sp., (Fig. 8C, E), but more protruding, perhaps because of a better preservation at this location. The only certitude is that it does not correspond to any other decorated bothremydid examined above by comparison with *Sindhochelys*. Although similar to *Sindhochelys* by the marked decoration and different from taphrosphyines and other examined bothremydid from France and India, the fragment remains undetermined.

DISCUSSION AND CONCLUSIONS

SYSTEMATIC AFFINITIES

Within Bothremyidae, *Sindhochelys ragei* n. gen., n. sp. appears as a new littoral form. At first instance, it seems to share the decoration and plastral scute pattern with Tahrosphyini. The study shows that these characters differ in the way they are made: among the species sharing the generalized scute plastral pattern and a marked decoration, *S. ragei* n. gen., n. sp. is unique in the details of these elements besides it is unique in the shape

of its carapace and the contour of the constituting elements. Thus, it is different from Taphrosphyini (*sensu nostro*), as well as from *Carteremys lethii* group from India and from *Elochelys perfecta* from France which are not closely related taxa.

Sindhochelys ragei n. gen., n. sp. cannot be referred to any of the few bothremydids known by their shell from Paleocene times: – either decorated: *Taphrosphys sulcatus*, *T. congolensis* (the other taphrosphyine Danian forms not being known by their shell [Gaffney *et al.* 2006]); – or not decorated: *Puentemys* from Colombia, the undefined form from Maria Farinha Fm. of Pernambuco, Brazil, DNPM (observed by Campos de Almeida & Broin [1981]) and “*Podocnemis*” *indica* Lydekker, 1885 from the Paleocene beds of Nila (Salt Range, Pakistan). In addition to *S. ragei* n. gen., n. sp., the latter is the only known other Paleocene Pakistani bothremydids, and both are closer in age. But this species is of an undetermined genus, known only by a drawing of a dorsal carapace, without ornamentation, carinated on the figure (although indicated as not carinated in Lydekker’s text, being perhaps just tectiform), narrow and oval, not at all similar to that of *S. ragei* n. gen., n. sp., and thus indicates no close relationship between the two. Moreover, *S. ragei* n. gen., n. sp. cannot be referred to any Cretaceous form defined by its shell and/or skull. In the absence of skull or shell in all the species, the phylogenetic affinities of *S. ragei* n. gen., n. sp. cannot yet be defined.

Large shells of bothremydids have been reported from several Maastrichtian stratigraphic intervals of the Mt Indamane – Mt In Tahout area of the Iullemeden basin (Greigert 1966; Moody & Sutcliffe 1991; Lapparent de Broin *et al.* 2020), Niger. The decoration and plastral pattern of these undescribed shells are unknown. *Ilatardia* Pérez-García, 2019b, was described by a skull alone in another Maastrichtian layer of the same basin, at Ilatarda. In absence of its shell, any comparison with *S. ragei* n. gen., n. sp. is possible. The skull of *Ilatardia* is very close to that of *Nigeremys* and we refer *Ilatardia* to the Nigeremydini (*sensu* Lapparent de Broin & Guntupalli 2020). Since the rare shells known (complete or fragmentary) in this tribe (Lapparent de Broin & Werner 1998; Moody & Sutcliffe 1991; O’Leary *et al.* 2019) are not decorated by raised well protruding polygons, it is probable that *Ilatardia* is not related to *S. ragei* n. gen., n. sp. (Lapparent de Broin *et al.* 2020). Other decorated forms existed in the Iullemeden basin, but they are undetermined and too poorly known to be fruitfully compared with *S. ragei* n. gen., n. sp. (see below).

PALEOGEOGRAPHIC CONSIDERATIONS

During the early Paleocene, the Ranikot area, as part of the Indian Subcontinent, was separated from Madagascar and from Africa, but closer to the Madagascan coasts than to the northern part of Africa where bothremydids are known at that time. The Bothremydidae are a Gondwanan taxon, the earliest representatives of which are from the early Cretaceous of northern Gondwana (Broin 1988). There are two main paleobiogeographic hypotheses to explain the presence of bothremydids in Pakistan. *Sindhochelys ragei* n. gen., n. sp. and “*Podocnemis indica*” may have evolved from forms which

inhabited eastern Gondwana before the Gondwana breakup between the Indian Subcontinent and Madagascar, during the early late Cretaceous, but their shells are unknown. Undecorated and decorated bothremydids are known in the Albian to Maastrichtian of India (Ayyasami & Das 1990; Lapparent de Broin & Guntupalli 2020) and Madagascar (Lapparent de Broin & Werner 1998; Lapparent de Broin 2000; Gaffney *et al.* 2006). *Carteremys lethii* from the Maastrichtian of India is the only decorated form but we demonstrated previously that it is not closely related to *S. ragei* n. gen., n. sp. An undetermined shell (preserved with some other poor elements in the MNHN.F collections) has been reported in the Cenomanian of Madagascar, but it is not decorated as in *S. ragei* n. gen., n. sp. (Lapparent de Broin & Werner 1998). Other forms are defined from the Cenomanian of Madagascar, but their shell is unknown (Gaffney *et al.* 2006).

Another hypothesis is that African bothremydids dispersed across the Neotethys to the Indian Subcontinent as it is attested for adapisoriculid mammals (Prasad *et al.* 2010). The age of this dispersal event(s) remains contentious, but it is worth noting that bothremydids are coastal turtles without the paddles of marine turtles, and their dispersion depends of favorable currents, and only between close areas and along the coasts. During the late Maastrichtian to early Paleocene, no bothremydids is known in the eastern part of Africa, but the family is well diversified in the northern part and in the surrounding Neotethysian areas, as early as Cenomanian. A current of northwestern-southeastern direction initiated in Africa or a relay between continents by a volcanic archipelago may have possibly favored the dispersal of littoral bothremydids during the late Cretaceous or the earliest Paleocene. Decorated bothremydids occurred in the Iullemeden basin (Niger), which was situated on the western border of the Trans-Saharan seaway and connected to the Neotethys. Besides other well-identified tribes including decorated taphrosphyines, and undecorated Nigeremydini and Bothremydini, Bothremydidae indet. are reported from the Maastrichtian Ammonite Hills Formation (Egypt) (Lapparent de Broin & Werner 1998; Lapparent de Broin 2000) but they are not decorated. In contrast, isolated plates (MNHN.F 1967.27 coll.) of undefined bothremydids from the Iullemeden of Niger, display a rather similar decoration to that of *S. ragei* n. gen., n. sp. These undefined bothremydids were recovered from the “papyraceous shales bed” of the Mt In Tahout area, which is considered Thanetian in age according to the oil-geologists who discovered the fossils (Lapparent de Broin *et al.* 2020), and they correspond to the Kao Member of the Garadoua Formation, which is exposed in the Iullemeden basin (Dikouma *et al.* 1994). Thus, they are probably younger than *S. ragei* n. gen., n. sp. from Ranikot. Another decorated peripheral of a bothremydids from Mt Indamane (of the same basin) is figured (Michaut 2002, 2007). This specimen referred as to “Testudines” by Michaut (2002) is about the size of *S. ragei* n. gen., n. sp., and its decoration with wide protruding polygons is similar. The specimen might come from the same Paleocene level as the elements of the neighboring Mt In Tahout level (Lapparent de Broin *et al.* 2020). Although these fossils are still

insufficient, their presence suggests that dispersal corridors might have occurred along the Trans-Saharan seaway, thus possibly connecting biogeographically Eastern Africa and India during the Maastrichtian or the Paleocene.

The occurrences of *Sindhochelys ragei* n. gen., n. sp. and “*Podocnemis*” *indica* indicate that the bothremydids were relatively diversified in the western margin of the Indian Subcontinent during the Paleocene. During the early Eocene, pleurodirans are represented by undetermined Pelomedusoides in India (Smith *et al.* 2016) and perhaps also in the Eocene of Pakistan (Broin 1987).

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